

From: Myers, Carla
Sent: Tuesday, November 14, 2006 7:29 AM
To: STIC-Biotech/ChemLib
Subject: sequence search for 10/823197

Please search:

- 1) SEQ ID NO: 7
- 2) please perform a score over length search of SEQ ID NO: 7, with a minimum length of 8 nucleotides and a maximum length of 50 nucleotides, and a score over length cutoff of 70%.

Please provide an alignment in SCORE of the first 50 results (there is no need to provide a printed version of the results).

Thank you-

Carla Myers
AU 1634
Remsen Bldg / Rm 2E79
Mailbox: REM 2C70
571-272-0747

17

ME

Searcher: _____
Searcher Phone: _____
Date Searcher Picked up: _____
Date completed: _____
Searcher Prep Time: _____
Online Time: _____

Type of Search
NA# _____ AA# _____
S/L: _____ Oligomer: _____
Encode/Transl: _____
Structure #: _____ Text: _____
Inventor: _____ Litigation: _____

Vendors and cost where applicable
STN: _____
DIALOG: _____
QUESTEL/ORBIS: _____
LEXIS/NEXIS: _____
SEQUENCE SYSTEM: _____
WWW/Internet: _____
Other (Specify): _____

Date completed: _____

Searcher: Beverly e 2528

Terminal time: _____

Elapsed time: _____

CPU time: _____

Total time: _____

Number of Searches: _____

Number of Databases: _____

Search Site

_____ STIC

_____ CM-1

_____ Pre-S

Type of Search

_____ N.A. Sequence

_____ A.A. Sequence

_____ Structure

_____ Bibliographic

Vendors

_____ IG

_____ STN

_____ Dialog

_____ APS

_____ Geninfo

_____ SDC

_____ DARC/Questel

✓ Other CGN

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Myers, C
10/823197
Seq:ID 7

GenCore version 5.1.9
Copyright (c) 1993 - 2006 Bioceleration Ltd.

OM nucleic - nucleic search, using sw model

Run on: November 21, 2006, 14:27:35 ; Search time 0.001 Seconds
(without alignment)
60.656 Million cell updates/sec

Title: US-10-823-197-7

Perfect score: 17

Sequence: 1 CTCAGCAACTCTCTAT 17

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 0.5

Searched: 156 seqs, 1784 residues

Total number of hits satisfying chosen parameters: 312

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 156 summaries

Database : rgedb:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	17	100.0	17	A84740	ACCESSION: A84740
3	17	100.0	17	AR144074	ACCESSION: AR144074
4	17	100.0	17	BD070852	ACCESSION: BD070852
5	17	100.0	17	BD085841	ACCESSION: BD085841
6	17	100.0	17	BD124538	ACCESSION: BD124538
7	17	100.0	17	BD246771	ACCESSION: BD246771
8	17	100.0	17	BD271249	ACCESSION: BD271249
9	17	100.0	17	CQ918145	ACCESSION: CQ918145
10	17	100.0	17	CS231184	ACCESSION: CS231184
11	17	100.0	17	AR282877	ACCESSION: AR282877
12	17	100.0	17	I85578	ACCESSION: I85578
13	17	100.0	17	AR487085	ACCESSION: AR487085
14	17	100.0	17	AR490513	ACCESSION: AR490513
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16	17	100.0	17	AR532946	ACCESSION: AR532946
17	17	100.0	17	AR534188	ACCESSION: AR534188
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DEFINITION A82477
ACCESSION A82477
VERSION A82477.1 GI:6732221
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
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unidentified
unclassified sequences.
Duff,G. and Cox,A.
PREDICTION OF INFLAMMATORY DISEASE ASSOCIATED WITH IL-1 GENELOC1
POLYMORPHISMS
Patent: WO 9854359-A 15 03-DEC-1998;
DUFF GORDON (GB); COX ANGELA (GB)
Location/Qualifiers
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DEFINITION Sequence 1 from Patent WO9844150.
ACCESSION A84740
VERSION A84740.1 GI:6733608
KEYWORDS
SOURCE
ORGANISM
unidentified
unclassified sequences.
REFERENCE 1 (bases 1 to 17)
AUTHORS Specter,T.D. and Keen,R.W.
TITLE POLYMORPHISMS OF AN IL-1 RECEPTOR ANTAGONIST GENE
JOURNAL Patent: WO 9844150-A 1 08-OCT-1998;
GEMINI RESEARCH LTD (GB); SPECTOR TIMOTHY DAVID (GB)
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Location/Qualifiers
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DEFINITION Sequence 1 from patent US 6210877.
ACCESSION AR144074
VERSION AR144074.1 GI:15105941
KEYWORDS
SOURCE
ORGANISM
Unknown.
Unclassified.
REFERENCE 1 (bases 1 to 17)
AUTHORS Francis,S.E.; Crossman,D.C. and Duff,G.W.
TITLE Prediction of coronary artery disease
JOURNAL Patent: US 6210877-A 1 03-APR-2001;
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DEFINITION Prediction of coronary artery disease.
ACCESSION  BD070852
VERSION    BD070852.1 GI:22616455
KEYWORDS   JP 2001514522-A/1.
SOURCE     synthetic construct
ORGANISM   other sequences; artificial sequences.
REFERENCE  1 (bases 1 to 17)
AUTHORS    Francis,S.E., Crossman,D.C. and Duff,G.W.
TITLE      Prediction of coronary artery disease
JOURNAL    Patent: JP 2001514522-A 1 11-SEP-2001;
           INTERLEUKIN GENETICS INC
COMMENT    OS Artificial Sequence
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DEFINITION Methods of diagnosing and treating chronic obstructive airway
ACCESSION  BD085841
VERSION    BD085841.1 GI:22631451
KEYWORDS   JP 2001522586-A/5.
SOURCE     unidentified
ORGANISM   unclassified sequences.
REFERENCE  1 (bases 1 to 17)
AUTHORS    Duff,G.W., Giovain,M., Barnes,P.J. and Rim,S.
TITLE      Methods of diagnosing and treating chronic obstructive airway
JOURNAL    Patent: JP 2001522586-A 5 20-NOV-2001;
           INTERLEUKIN GENETICS INC
COMMENT    OS Unidentified
           PN JP 2001522586-A/5
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ACCESSION  BD246771
VERSION    BD246771.1 GI:33056541
KEYWORDS   JP 2002533096-A/1.
SOURCE     synthetic construct
ORGANISM   other sequences; artificial sequences.
REFERENCE  1 (bases 1 to 17)
AUTHORS    Giovine,F.S.D. and Duff,G.W.
TITLE      Diagnostics and therapeutics for sepsis
JOURNAL    Patent: JP 2002533096-A 1 08-OCT-2002;

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DEFINITION Prediction of inflammatory disease associated with IL-1 geneloci
ACCESSION  BD124538
VERSION    BD124538.1 GI:23219483
KEYWORDS   JP 2002500513-A/15.
SOURCE     unidentified
ORGANISM   unclassified sequences.
REFERENCE  1 (bases 1 to 17)
AUTHORS    Duff,G., Cox,A., Camp,N.J. and Giovine,F.S.D.
TITLE      Prediction of inflammatory disease associated with IL-1 geneloci
JOURNAL    Patent: JP 2002500513-A 15 08-JAN-2002;
           INTERLEUKIN GENETICS INC
COMMENT    OS Unidentified
           PN JP 2002500513-A/15
           PD 08-JAN-2002
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           PR 29-MAY-1997 GB 9711040.7
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RESULT 7
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DEFINITION Diagnostics and therapeutics for sepsis.
ACCESSION  BD246771
VERSION    BD246771.1 GI:33056541
KEYWORDS   JP 2002533096-A/1.
SOURCE     synthetic construct
ORGANISM   other sequences; artificial sequences.
REFERENCE  1 (bases 1 to 17)
AUTHORS    Giovine,F.S.D. and Duff,G.W.
TITLE      Diagnostics and therapeutics for sepsis
JOURNAL    Patent: JP 2002533096-A 1 08-OCT-2002;

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INTERLEUKIN GENETICS INC
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PN JP 2002533096-A/1
PD 08-OCT-2002
PF 01-NOV-1999 JP 2000599732
PR 30-OCT-1998 US 09/183850
PI FRANCESCO S DI GIOVINE, GORDON W DUFF
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DEFINITION Prediction of risk of interstitial lung disease.
ACCESSION BD271249
VERSION BD271249.1 GI:33081017
KEYWORDS JP 2002540801-A/21.
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1 (bases 1 to 17)
AUTHORS Duff,G.W., Giovine,F.S.D. and Whyte,M.
TITLE Prediction of risk of interstitial lung disease
JOURNAL Patent: JP 2002540801-A 21 03-DEC-2002;
INTERLEUKIN GENETICS INC
COMMENT OS Artificial Sequence
PN JP 2002540801-A/21
PD 03-DEC-2002
PF 31-MAR-2000 JP 2000609606
PR 02-APR-1999 US 09/286108
PI GORDON W DUFF,FRANCESCO SAVERIO DI GIOVINE,MORIA WHITE PC
C12N15/09,A61K31/57,A61K31/7088,A61K38/00,A61K45/00,A61K48/00, PC
A61P11/00,
PC C12Q1/02,C12Q1/68,G01N33/15,G01N33/50,G01N33/53,G01N33/566, PC
C12N15/00,
PC A61K37/02
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FH Key Location/Qualifiers
FT source
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/organism='Artificial Sequence'.
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Query Match 100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 8;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

INTERLEUKIN GENETICS INC
OS Artificial Sequence
PN JP 2002533096-A/1
PD 08-OCT-2002
PF 01-NOV-1999 JP 2000599732
PR 30-OCT-1998 US 09/183850
PI FRANCESCO S DI GIOVINE, GORDON W DUFF
PC C12N15/09,A61K38/00,A61K45/00,A61P31/00,C12Q1/68,C12N15/00, PC
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CC Description of Artificial Sequence: primer
FH Key Location/Qualifiers
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Query Match 100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 8;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 9
CQ918145
LOCUS CQ918145 17 bp DNA linear PAT 23-NOV-2004
DEFINITION Sequence 29 from Patent WO2004097045.
ACCESSION CQ918145
VERSION CQ918145.1 GI:56208292
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1
AUTHORS Brown,M.A.
TITLE Diagnostic assay for ankylosing spondylitis
JOURNAL Patent: WO 2004097045-A 29 11-NOV-2004;
ISIS INNOVATION LIMITED (GB)
FEATURES Location/Qualifiers
source 1..17
/organism='synthetic construct'
/mol_type='unassigned DNA'
/db_xref='taxon:32630'
/notes='Primer'

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Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 10
CS231184
LOCUS CS231184 17 bp DNA linear PAT 15-DEC-2005
DEFINITION Sequence 22 from Patent WO2005108619.
ACCESSION CS231184
VERSION CS231184.1 GI:83698459
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1
AUTHORS Duff,G.W.
TITLE Diagnostics and therapeutics for diseases associated with an
il-1inflammatory haplotype
JOURNAL Patent: WO 2005108619-A 22 17-NOV-2005;
Interleukin Genetics, Inc. (US)
FEATURES Location/Qualifiers
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/mol_type='unassigned DNA'
/db_xref='taxon:32630'
/notes='Description of Artificial Sequence: chemically
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Best Local Similarity 100.0%; Pred. No. 8;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 11
AR282877
LOCUS AR282877 17 bp DNA linear PAT 10-APR-2003
DEFINITION Sequence 1 from patent US 6524795.
ACCESSION AR282877
VERSION AR282877.1 GI:29719679
KEYWORDS Unknown.
SOURCE Unknown.

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Unclassified.
 1 (bases 1 to 17)
 Francis, S.E., Crossman, D.C., Duff, G.W., Kornman, K.S. and
 Stephenson, K.
 TITLE
 Journal
 Patent: US 6524795-A 1 25-FEB-2003;
 Location/Qualifiers
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 /mol_type="genomic DNA"

Query Match 100.0%; Score 17; DB 1; Length 17;
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Qy 1 CTCAGCAACTCTCTAT 17
 Db 1 CTCAGCAACTCTCTAT 17

RESULT 12
 I85578
 LOCUS I85578 17 bp DNA linear PAT 10-JUN-1998
 DEFINITION Sequence 1 from patent US 5698399.
 ACCESSION I85578
 VERSION I85578.1 GI:3205296
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE
 1 (bases 1 to 17)
 Duff, G.W., Russell, G. and Eastell, R.
 TITLE
 Detecting genetic predisposition for osteoporosis
 JOURNAL
 Patent: US 5698399-A 1 16-DEC-1997;
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 Db 1 CTCAGCAACTCTCTAT 17

RESULT 13
 AR487085
 LOCUS AR487085 17 bp DNA linear PAT 14-MAY-2004
 DEFINITION Sequence 22 from patent US 6706478.
 ACCESSION AR487085
 VERSION AR487085.1 GI:47252036
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE
 1 (bases 1 to 17)
 Duff, G.W., Cox, A., Camp, N.J. and di Giovine, F.S.
 TITLE
 Diagnostics and therapeutics for diseases associated with an IL-1
 inflammatory haplotype
 JOURNAL
 Patent: US 6706478-A 22 16-MAR-2004;
 Interleukin Genetics, Inc.; Waltham, MA;
 GBX;
 Location/Qualifiers
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Qy 1 CTCAGCAACTCTCTAT 17
 Db 1 CTCAGCAACTCTCTAT 17

RESULT 14
 AR490513
 LOCUS AR490513 17 bp DNA linear PAT 15-MAY-2004
 DEFINITION Sequence 5 from patent US 6713253.
 ACCESSION AR490513
 VERSION AR490513.1 GI:47257894
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE
 1 (bases 1 to 17)
 Duff, G.W., Richardson, P.R.S. and Rennie, I.G.
 TITLE
 Detecting genetic predisposition to sight-threatening diabetic
 retinopathy
 JOURNAL
 Patent: US 6713253-A 5 30-MAR-2004;
 Interleukin Genetics, Inc.; Waltham, MA;
 GBX;
 Location/Qualifiers
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Qy 1 CTCAGCAACTCTCTAT 17
 Db 1 CTCAGCAACTCTCTAT 17

RESULT 15
 AR493814
 LOCUS AR493814 17 bp DNA linear PAT 15-MAY-2004
 DEFINITION Sequence 7 from patent US 6720141.
 ACCESSION AR493814
 VERSION AR493814.1 GI:47266264
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE
 1 (bases 1 to 17)
 Crossman, D.C., Duff, G.W., Francis, S.E., Kornman, K.S. and
 Stephenson, K.
 TITLE
 Diagnostics and therapeutics for restenosis
 JOURNAL
 Patent: US 6720141-A 7 13-APR-2004;
 Interleukin Genetics, Inc.; Waltham, MA
 Location/Qualifiers
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Qy 1 CTCAGCAACTCTCTAT 17
 Db 1 CTCAGCAACTCTCTAT 17

RESULT 16
 AR532946
 LOCUS AR532946 17 bp DNA linear PAT 08-OCT-2004
 DEFINITION Sequence 9 from patent US 6730476.
 ACCESSION AR532946
 VERSION AR532946.1 GI:53922498


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AX067286
LOCUS AX067286 17 bp DNA linear PAT 24-JAN-2001
DEFINITION Sequence 22 from Patent WO0100880.
ACCESSION AX067286
VERSION AX067286.1 GI:12544910
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE
AUTHORS Duff,G.W., Cox,A., Camp,N.J. and di Giovine,F.S.
TITLE Diagnostics and therapeutics for diseases associated with an il-1
inflammatory haplotype
JOURNAL Patent: WO 0100880-A 22 04-JAN-2001;
Interleukin Genetics, Inc. (US)
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/db_xref="taxon:32630"
/note="primer"
Query Match 100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 8;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACTCCTAT 17
Db 1 CTCAGCAACTCCTAT 17

RESULT 22
AX360017
LOCUS AX360017 17 bp DNA linear PAT 13-FEB-2002
DEFINITION Sequence 3 from Patent WO200933.
ACCESSION AX360017
VERSION AX360017.1 GI:18675643
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.
REFERENCE
AUTHORS Duff,G.W. and Kornman,K.S.
TITLE Screening assays for identifying modulators of the inflammatory or
immune responses
JOURNAL Patent: WO 0200933-A 3 03-JAN-2002;
Interleukin Genetics, Inc. (US)
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/note="Primer"
Query Match 100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 8;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACTCCTAT 17
Db 1 CTCAGCAACTCCTAT 17

RESULT 23
AR045302/c
LOCUS AR045302/c 15 bp DNA linear PAT 29-SEP-1999
DEFINITION Sequence 95 from patent US 5817796.
ACCESSION AR045302
VERSION AR045302.1 GI:5966767
KEYWORDS Unknown.
SOURCE Unknown.
ORGANISM Unclassified.

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REFERENCE 1 (bases 1 to 15)
AUTHORS Stinchcomb,D.T., Draper,K., McSwiggen,J. and Jarvis,T.
TITLE C-myb ribozymes having 2'-5'-linked adenylate residues
JOURNAL Patent: US 5817796-A 95 06-OCT-1998;
FEATURES
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/organism="unknown"
/mol_type="unassigned DNA"
Query Match 63.5%; Score 10.8; DB 1; Length 15;
Best Local Similarity 85.7%; Pred. No. 47;
Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CTCAGCAACTCC 14
Db 14 CTCAGCAACTTC 1

RESULT 24
152354/c
LOCUS 152354 15 bp DNA linear PAT 07-OCT-1997
DEFINITION Sequence 95 from patent US 5646042.
ACCESSION 152354
VERSION 152354.1 GI:2473555
KEYWORDS Unknown.
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 15)
AUTHORS Stinchcomb,D.T., Draper,K., McSwiggen,J. and Jarvis,T.
TITLE C-myb targeted ribozymes
JOURNAL Patent: US 5646042-A 95 08-JUL-1997;
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/organism="unknown"
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Best Local Similarity 85.7%; Pred. No. 47;
Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CTCAGCAACTCC 14
Db 14 CTCAGCAACTTC 1

RESULT 25
AX587048/c
LOCUS AX587048 15 bp DNA linear PAT 10-JAN-2003
DEFINITION Sequence 70 from Patent WO02072883.
ACCESSION AX587048
VERSION AX587048.1 GI:27655923
KEYWORDS Saccharomyces cerevisiae (baker's yeast)
SOURCE Saccharomyces cerevisiae
ORGANISM Saccharomyces cerevisiae
REFERENCE 1
AUTHORS Roetger,A.
TITLE Nucleotide carrier for diagnosing and treating oral diseases
JOURNAL Patent: WO 02072883-A 70 19-SEP-2002;
Roetger, Antje (DE)
FEATURES
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/organism="Saccharomyces cerevisiae"
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/db_xref="taxon:4932"
Query Match 63.5%; Score 10.8; DB 1; Length 15;
Best Local Similarity 85.7%; Pred. No. 47;
Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 4 AGCAACTCCTAT 17

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RESULT 30
AR301565/c
LOCUS AR301565 11 bp DNA linear PAT 12-JUN-2003
DEFINITION Sequence 146 from patent US 6538173.
ACCESSION AR301565
VERSION AR301565.1 GI:31689367
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE
Unclassified.
AUTHORS Heber-Katz, E.
TITLE Compositions and methods for wound healing
JOURNAL Patent: US 6538173-A 146 25-MAR-2003;
The Wistar Institute; Philadelphia, PA;
WOK;
FEATURES
source
1..11
/mol_type="genomic DNA"
Query Match 52.9%; Score 9; DB 1; Length 11;
Best Local Similarity 100.0%; Pred. No. 42;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTCAGCAAC 9
Db 9 CTCAGCAAC 1
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RESULT 31
AR322159
LOCUS AR322159 12 bp DNA linear PAT 17-AUG-2003
DEFINITION Sequence 10 from patent US 6566061.
ACCESSION AR322159
VERSION AR322159.1 GI:33707703
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE
Unclassified.
AUTHORS Philibert, R.A., Gimms, E.I. and Delisi, L.
TITLE Identification of polymorphisms in the PCTG4 region of Xq13
JOURNAL Patent: US 6566061-A 10 20-MAY-2003;
The University of Iowa, as represented by the University of Iowa
Research Foundation and The United States of America as represented
by the Department of Health and Human Services; Iowa City, IA
FEATURES
source
1..12
/mol_type="genomic DNA"
Query Match 52.9%; Score 9; DB 1; Length 12;
Best Local Similarity 100.0%; Pred. No. 51;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 3 CAGCAACAC 11
Db 1 CAGCAACAC 9
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RESULT 32
BD239508
LOCUS BD239508 10 bp DNA linear PAT 17-JUL-2003
DEFINITION Preparation and use of superior vaccines.
ACCESSION BD239508
VERSION BD239508.1 GI:33049278
KEYWORDS JP 2002534056-A/926.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominiidae; Homo.

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REFERENCE
1 (bases 1 to 10)
AUTHORS Roberts, B.L. and Shankara, S.
TITLE Preparation and use of superior vaccines
JOURNAL Patent: JP 2002534056-A 926 15-OCT-2002;
GENZYME CORP
COMMENT
OS Homo sapiens (human)
PN JP 2002534056-A/926
PD 15-OCT-2002
PF 18-JUN-1999 JP 2000554749
PR 19-JUN-1998 US 60/090039, 19-JUN-1998 US 60/090040 PR
19-JUN-1998 US 60/090041, 19-JUN-1998 US 60/089853 PR
19-JUN-1998 US 60/089997, 19-JUN-1998 US 60/090079 PR
19-JUN-1998 US 60/090035, 19-JUN-1998 US 60/089993 PR
19-JUN-1998 US 60/089992, 19-JUN-1998 US 60/090072 PR
19-JUN-1998 US 60/089878, 19-JUN-1998 US 60/089991 PR
19-JUN-1998 US 60/090000, 19-JUN-1998 US 60/090043 PR
19-JUN-1998 US 60/089999, 19-JUN-1998 US 60/090048 PR
19-JUN-1998 US 60/090042, 19-JUN-1998 US 60/090036 PR
19-JUN-1998 US 60/090044, 19-JUN-1998 US 60/089844 PR
19-JUN-1998 US 60/090080, 19-JUN-1998 US 60/089833 PR
19-JUN-1998 US 60/089994, 19-JUN-1998 US 60/090077 PR
19-JUN-1998 US 60/090078, 19-JUN-1998 US 60/090047 PR
19-JUN-1998 US 60/090076, 19-JUN-1998 US 60/090045 PR
08-DEC-1998 US 60/111715
PI BRUCE L ROBERTS, SRINIVAS SHANKARA
PC C12N15/09, C12N15/09, A61K39/00, A61P35/00, A61P37/04, C12N1/15, PC
C12N1/19,
PC C12N1/21, C12N5/10, G01N33/15, G01N33/50, G01N33/53, G01N33/566, PC
G01N37/00,
PC C12N15/00, C12N5/00, C12N15/00
CC Preparation and use of superior vaccines
FH Key Location/Qualifiers
FT source 1..10
/organism="Homo sapiens (human)"
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Location/Qualifiers
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Best Local Similarity 90.0%; Pred. No. 41;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 CTCAGCAACA 10
Db 1 CTCAGCAAAA 10
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RESULT 33
BD239610/c
LOCUS BD239610 10 bp DNA linear PAT 17-JUL-2003
DEFINITION Preparation and use of superior vaccines.
ACCESSION BD239610
VERSION BD239610.1 GI:33049380
KEYWORDS JP 2002534056-A/1028.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominiidae; Homo.
REFERENCE
1 (bases 1 to 10)
AUTHORS Roberts, B.L. and Shankara, S.
TITLE Preparation and use of superior vaccines
JOURNAL Patent: JP 2002534056-A 1028 15-OCT-2002;
GENZYME CORP
COMMENT
OS Homo sapiens (human)
PN JP 2002534056-A/1028
PD 15-OCT-2002
PF 18-JUN-1999 JP 2000554749
PR 19-JUN-1998 US 60/090039, 19-JUN-1998 US 60/090040 PR
19-JUN-1998 US 60/090041, 19-JUN-1998 US 60/089853 PR
19-JUN-1998 US 60/089997, 19-JUN-1998 US 60/090079 PR

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19-JUN-1998 US 60/090035,19-JUN-1998 US 60/089993 PR
19-JUN-1998 US 60/089992,19-JUN-1998 US 60/090072 PR
19-JUN-1998 US 60/089878,19-JUN-1998 US 60/089991 PR
19-JUN-1998 US 60/090000,19-JUN-1998 US 60/090048 PR
19-JUN-1998 US 60/089999,19-JUN-1998 US 60/090043 PR
19-JUN-1998 US 60/090042,19-JUN-1998 US 60/090036 PR
19-JUN-1998 US 60/090044,19-JUN-1998 US 60/089844 PR
19-JUN-1998 US 60/090080,19-JUN-1998 US 60/089833 PR
19-JUN-1998 US 60/089994,19-JUN-1998 US 60/090077 PR
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19-JUN-1998 US 60/090076,19-JUN-1998 US 60/090045 PR
08-DEC-1998 US 60/111715
PI BRUCE L ROBERTS, SRINIVAS SHANKARA
PC C12N15/09,C12N15/09,A61K39/00,A61P35/00,A61P37/04,C12N1/15, PC
C12N1/19,
PC C12N1/21,C12N5/10,G01N33/15,G01N33/50,G01N33/53,G01N33/566, PC
G01N37/00,
PC C12N15/00,C12N5/00,C12N15/00
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FT Location/Qualifiers
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/db_xref="taxon:9606"

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Best Local Similarity 90.0%; Pred. No. 41;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

* Qy 1 CTCAGCAACA 10
Db 10 CTCAGCAACA 1

RESULT 34
CQ944980
LOCUS
DEFINITION Sequence 127 from Patent WO2004099445.
ACCESSION CQ944980
VERSION CQ944980.1 GI:56294321
KEYWORDS
ORGANISM synthetic construct
SOURCE synthetic construct
LOCATION/Qualifiers
other sequences; artificial sequences.
REFERENCE 1
AUTHORS Kahl,G., Winter,P., Krueger,D., Reich,S., Matsumura,H. and Terauchi,R.
TITLE Use of a type iii restriction enzyme to isolate identification tags comprising more than 25 nucleotides
JOURNAL Patent: WO 2004099445-A 127 18-NOV-2004; Iwate Prefectural Government (JP)
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/db_xref="taxon:32630"
/notes="Description of Artificial Sequence:Synthetic DNA (Tag Sequence)"

Query Match 49.4%; Score 8.4; DB 1; Length 10;
Best Local Similarity 90.0%; Pred. No. 41;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 3 CAGCAACT 12
Db 1 CAGCAACT 10

RESULT 35
AX153002
LOCUS
DEFINITION Sequence 917 from Patent WO0138577.
ACCESSION AX153002
VERSION AX153002.1 GI:14534653
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Velculescu,V.E., Vogelstein,B. and Kinzler,K.W.
TITLE Human transcriptomes
JOURNAL Patent: WO 0138577-A 917 31-MAY-2001; The Johns Hopkins University (US)
FEATURES
source 1..10
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/mol_type="unassigned DNA"
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Query Match 49.4%; Score 8.4; DB 1; Length 10;
Best Local Similarity 90.0%; Pred. No. 41;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 6 CAACACTCCT 15
Db 1 CAACACTCCT 10

RESULT 36
AX377359
LOCUS
DEFINITION Sequence 23 from Patent WO212499.
ACCESSION AX377359
VERSION AX377359.1 GI:19573645
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Kliem,S.B., Koshiy,B. and Lanz,E.M.
TITLE Haplotypes of the ntfs gene
JOURNAL Patent: WO 0212499-A 23 14-FEB-2002; Genessee Pharmaceuticals, Inc. (US)
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Best Local Similarity 90.0%; Pred. No. 41;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 5 GCAACACTCC 14
Db 1 GCAACACTCC 10

RESULT 37
CQ832726/c
LOCUS
DEFINITION Sequence 97 from Patent WO2004059002.
ACCESSION CQ832726
VERSION CQ832726.1 GI:50832333
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;

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REFERENCE
AUTHORS      Petersohn,D., Schlottmann,K., Gassenmeier,T., Holtkoetter,O.,
              Conrad,M. and Hofmann,K.
TITLE        Method for determining the homeostasis of hairy skin
JOURNAL      Patent: WO 2004059002-A 97 15-JUL-2004;
              Henkel Kommanditgesellschaft auf Aktien (DE)
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Query Match  49.4%; Score 8.4; DB 1; Length 11;
Best Local Similarity 90.0%; Pred. No. 51;
Matches      9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy           1 CTCAGCAACA 10
Db           11 CCCAGCAACA 2

RESULT 38
LOCUS      CQ833174
DEFINITION Sequence 545 from Patent WO2004059002.
ACCESSION  CQ833174
VERSION     CQ833174.1 GI:50832781
KEYWORDS   Homo sapiens (human)
SOURCE     Homo sapiens
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
            Hominidae; Homo.
REFERENCE  1
AUTHORS    Petersohn,D., Schlottmann,K., Gassenmeier,T., Holtkoetter,O.,
            Conrad,M. and Hofmann,K.
TITLE      Method for determining the homeostasis of hairy skin
JOURNAL    Patent: WO 2004059002-A 545 15-JUL-2004;
            Henkel Kommanditgesellschaft auf Aktien (DE)
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Qy           3 CAGCAACT 12
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RESULT 39
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DEFINITION Sequence 256 from Patent WO2004059001.
ACCESSION  CQ835198
VERSION     CQ835198.1 GI:50834732
KEYWORDS   Homo sapiens (human)
SOURCE     Homo sapiens
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
            Hominidae; Homo.
REFERENCE  1
AUTHORS    Petersohn,D., Schlottmann,K., Gassenmeier,T., Holtkoetter,O.,
            Conrad,M. and Hofmann,K.
TITLE      Method for determining markers of human facial skin
JOURNAL    Patent: WO 2004059001-A 256 15-JUL-2004;
            Henkel Kommanditgesellschaft auf Aktien (DE)
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LOCUS      CS058107
DEFINITION Sequence 4 from Patent WO2005028671.
ACCESSION  CS058107
VERSION     CS058107.1 GI:62551059
KEYWORDS   Homo sapiens (human)
SOURCE     Homo sapiens
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
            Hominidae; Homo.
REFERENCE  1
AUTHORS    Holtkoetter,O., Petersohn,D., Schlottmann,K., Giesen,M. and
            Kessler-Becker,D.
TITLE      Method for determining hair cycle markers
JOURNAL    Patent: WO 2005028671-A 4 31-MAR-2005;
            Henkel Kommanditgesellschaft auf Aktien (DE)
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Qy           6 CAACACTCCT 15
Db           1 CAACACTCCT 10

RESULT 41
LOCUS      CS058332
DEFINITION Sequence 229 from Patent WO2005028671.
ACCESSION  CS058332
VERSION     CS058332.1 GI:62551515
KEYWORDS   Homo sapiens (human)
SOURCE     Homo sapiens
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
            Hominidae; Homo.
REFERENCE  1
AUTHORS    Holtkoetter,O., Petersohn,D., Schlottmann,K., Giesen,M. and
            Kessler-Becker,D.
TITLE      Method for determining hair cycle markers
JOURNAL    Patent: WO 2005028671-A 229 31-MAR-2005;
            Henkel Kommanditgesellschaft auf Aktien (DE)
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Qy 6 CAACACTCCT 15
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RESULT 42
AX623499
LOCUS AX623499 11 bp DNA linear PAT 21-FEB-2003
DEFINITION Sequence 540 from Patent WO02053774.
ACCESSION AX623499
VERSION AX623499.1 GI:28451440
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM

REFERENCE
AUTHORS Petersohn,D., Conradt,M. and Hofmann,K.
TITLE Method for determining homeostasis of the skin
JOURNAL Patent: WO 02053774-A 540 11-JUL-2002;
Henkel Kommanditgesellschaft auf Aktien (DE)
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Qy 5 GCAACACTCC 14
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RESULT 43
AX624606
LOCUS AX624606 11 bp DNA linear PAT 21-FEB-2003
DEFINITION Sequence 1647 from Patent WO02053774.
ACCESSION AX624606
VERSION AX624606.1 GI:28452547
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM

REFERENCE
AUTHORS Petersohn,D., Conradt,M. and Hofmann,K.
TITLE Method for determining homeostasis of the skin
JOURNAL Patent: WO 02053774-A 1647 11-JUL-2002;
Henkel Kommanditgesellschaft auf Aktien (DE)
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Qy 7 AACACTCCTA 16
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Db 1 AACATTCCTA 10

RESULT 44
AX627166
LOCUS AX627166 11 bp DNA linear PAT 21-FEB-2003
DEFINITION Sequence 4207 from Patent WO02053774.
ACCESSION AX627166
VERSION AX627166.1 GI:28455204
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM

Query Match 49.4%; Score 8.4; DB 1; Length 11;
Best Local Similarity 90.0%; Pred. No. 51;
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AX624745
LOCUS AX624745 11 bp DNA linear PAT 21-FEB-2003
DEFINITION Sequence 1786 from Patent WO02053774.
ACCESSION AX624745
VERSION AX624745.1 GI:28452686
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM

REFERENCE
AUTHORS Petersohn,D., Conradt,M. and Hofmann,K.
TITLE Method for determining homeostasis of the skin
JOURNAL Patent: WO 02053774-A 1786 11-JUL-2002;
Henkel Kommanditgesellschaft auf Aktien (DE)
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Query Match 49.4%; Score 8.4; DB 1; Length 11;
Best Local Similarity 90.0%; Pred. No. 51;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 4 AGCAACACTC 13
| | | | |
Db 1 AGAAACACTC 10

RESULT 45
AX626573
LOCUS AX626573 11 bp DNA linear PAT 21-FEB-2003
DEFINITION Sequence 3614 from Patent WO02053774.
ACCESSION AX626573
VERSION AX626573.1 GI:28454611
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM

REFERENCE
AUTHORS Petersohn,D., Conradt,M. and Hofmann,K.
TITLE Method for determining homeostasis of the skin
JOURNAL Patent: WO 02053774-A 3614 11-JUL-2002;
Henkel Kommanditgesellschaft auf Aktien (DE)
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Best Local Similarity 90.0%; Pred. No. 51;
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Qy 6 CAACACTCCT 15
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Db 1 CAACATTCCT 10

RESULT 46
AX627166
LOCUS AX627166 11 bp DNA linear PAT 21-FEB-2003
DEFINITION Sequence 4207 from Patent WO02053774.
ACCESSION AX627166
VERSION AX627166.1 GI:28455204
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM

REFERENCE
AUTHORS Petersohn,D., Conradt,M. and Hofmann,K.
TITLE Method for determining homeostasis of the skin
JOURNAL Patent: WO 02053774-A 3614 11-JUL-2002;
Henkel Kommanditgesellschaft auf Aktien (DE)
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Best Local Similarity 90.0%; Pred. No. 51;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 6 CAACACTCCT 15
| | | | |
Db 1 CAACATTCCT 10

RESULT 47
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LOCUS AX627166 11 bp DNA linear PAT 21-FEB-2003
DEFINITION Sequence 4207 from Patent WO02053774.
ACCESSION AX627166
VERSION AX627166.1 GI:28455204
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM

REFERENCE
AUTHORS Petersohn,D., Conradt,M. and Hofmann,K.
TITLE Method for determining homeostasis of the skin
JOURNAL Patent: WO 02053774-A 3614 11-JUL-2002;
Henkel Kommanditgesellschaft auf Aktien (DE)
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Query Match 49.4%; Score 8.4; DB 1; Length 11;
Best Local Similarity 90.0%; Pred. No. 51;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.

REFERENCE

AUTHORS Petersohn,D., Conrad,M. and Hofmann,K.
TITLE Method for determining homeostasis of the skin
JOURNAL Patent: WO 02053774-A 4207 11-JUL-2002;
Henkel Kommanditgesellschaft auf Aktien (DE)

FEATURES

Location/Qualifiers
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/mol_type="unassigned DNA"
/db_xref="taxon:9606"

Query Match 49.4%; Score 8.4; DB 1; Length 11;
Best Local Similarity 90.0%; Pred. No. 51;
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QY

6 CAACACTCCT 15

1 CAGCACTCCT 10

RESULT 47
AX630920
LOCUS AX630920 11 bp DNA linear PAT 21-FEB-2003
DEFINITION Sequence 7961 from Patent WO02053774.
ACCESSION AX630920
VERSION AX630920.1 GI:28458960
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.
1
Petersohn,D., Conrad,M. and Hofmann,K.
TITLE Method for determining homeostasis of the skin
JOURNAL Patent: WO 02053774-A 7961 11-JUL-2002;
Henkel Kommanditgesellschaft auf Aktien (DE)

FEATURES

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QY

5 GCAACACTCC 14

1 GCAAGACTCC 10

RESULT 48
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LOCUS AX632027 11 bp DNA linear PAT 21-FEB-2003
DEFINITION Sequence 9069 from Patent WO02053774.
ACCESSION AX632027
VERSION AX632027.1 GI:28467642
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.
1
Petersohn,D., Conrad,M. and Hofmann,K.
TITLE Method for determining homeostasis of the skin
JOURNAL Patent: WO 02053774-A 9069 11-JUL-2002;
Henkel Kommanditgesellschaft auf Aktien (DE)

FEATURES

Location/Qualifiers
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/organism="Homo sapiens"
/mol_type="unassigned DNA"
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Query Match 49.4%; Score 8.4; DB 1; Length 11;
Best Local Similarity 90.0%; Pred. No. 51;
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QY

7 AACACTCCTA 16

1 AACATTCTTA 10

RESULT 49
AX632166
LOCUS AX632166 11 bp DNA linear PAT 21-FEB-2003
DEFINITION Sequence 9208 from Patent WO02053774.
ACCESSION AX632166
VERSION AX632166.1 GI:28467781
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.
1
Petersohn,D., Conrad,M. and Hofmann,K.
TITLE Method for determining homeostasis of the skin
JOURNAL Patent: WO 02053774-A 9208 11-JUL-2002;
Henkel Kommanditgesellschaft auf Aktien (DE)

FEATURES

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Best Local Similarity 90.0%; Pred. No. 51;
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QY

4 AGCAACACTC 13

1 AGAAACACTC 10

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LOCUS AX708088 11 bp DNA linear PAT 04-APR-2003
DEFINITION Sequence 24 from Patent WO03014387.
ACCESSION AX708088
VERSION AX708088.1 GI:29564039
KEYWORDS synthetic construct
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.

REFERENCE
AUTHORS Wojnowski,L. and Presecan-Siedel,E.
TITLE Polymorphisms in the human gene for cypla2 and their use in
diagnostic and therapeutic applications
JOURNAL Patent: WO 03014387-A 24 20-FEB-2003;
Epidauros Biotechnologie AG (DE)
FEATURES
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1. .11
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QY

8 ACACCTCCTAT 17

11

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c 114	9.4	55.3	12	1	ABH83410	Oligonucleotide pr	c 187	9.4	55.3	13	1	ABC86466	Oligonucleotide SE
c 115	9.4	55.3	12	1	ABH87181	Oligonucleotide pr	c 188	9.4	55.3	13	1	ABF32704	Oligonucleotide SE
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121	9.4	55.3	12	1	ABI15864	Oligonucleotide pr	c 194	9.4	55.3	13	1	ABF73020	Oligonucleotide SE
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c 124	9.4	55.3	12	1	ABI79343	Oligonucleotide pr	c 197	9.4	55.3	13	1	ABC96629	Oligonucleotide SE
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c 126	9.4	55.3	13	1	ABF69369	Oligonucleotide SE	c 199	9.4	55.3	13	1	ABC78901	Oligonucleotide SE
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c 140	9.4	55.3	13	1	ABC24578	Oligonucleotide SE	c 213	9.4	55.3	13	1	ABF27389	Oligonucleotide SE
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c 148	9.4	55.3	13	1	ABF08236	Oligonucleotide SE	c 221	9	52.9	10	1	AAF41735	Yeast NORF gene SA
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c 150	9.4	55.3	13	1	ABF27388	Oligonucleotide SE	c 223	9	52.9	11	1	AZ18836	Murine C57BL/6 SAG
c 151	9.4	55.3	13	1	ABF53293	Oligonucleotide SE	c 224	9	52.9	12	1	AZ52017	B-cell mRNA ribozyme
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c 166	9.4	55.3	13	1	ABF53999	Oligonucleotide SE	c 239	8	51.8	12	1	ABH69162	Oligonucleotide pr
c 167	9.4	55.3	13	1	ABH59111	Oligonucleotide SE	c 240	8	51.8	12	1	ABH22817	Oligonucleotide pr
c 168	9.4	55.3	13	1	ABH71081	Oligonucleotide SE	c 241	8	51.8	12	1	ABH43805	Oligonucleotide pr
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c 170	9.4	55.3	13	1	ABC86465	Oligonucleotide SE	c 243	8	51.8	12	1	ABH103049	Oligonucleotide pr
c 171	9.4	55.3	13	1	ABH27109	Oligonucleotide SE	c 244	8	51.8	12	1	ABH32041	Oligonucleotide pr
c 172	9.4	55.3	13	1	ABF53292	Oligonucleotide SE	c 245	8	51.8	12	1	ABH82739	Oligonucleotide pr
c 173	9.4	55.3	13	1	ABH07983	Oligonucleotide SE	c 246	8	51.8	12	1	ABH14114	Oligonucleotide pr
c 174	9.4	55.3	13	1	ABC86467	Oligonucleotide SE	c 247	8	51.8	12	1	ABH70127	Oligonucleotide pr
c 175	9.4	55.3	13	1	ABF78085	Oligonucleotide SE	c 248	8	51.8	12	1	ABH72016	Oligonucleotide pr
c 176	9.4	55.3	13	1	ABC00319	Oligonucleotide SE	c 249	8	51.8	12	1	ABH90061	Oligonucleotide pr
c 177	9.4	55.3	13	1	ABF90353	Oligonucleotide SE	c 250	8	51.8	12	1	ABH89055	Oligonucleotide pr
c 178	9.4	55.3	13	1	ABH71075	Oligonucleotide SE	c 251	8	51.8	12	1	ABH53259	Oligonucleotide pr
c 179	9.4	55.3	13	1	ABH71084	Oligonucleotide SE	c 252	8	51.8	12	1	ABH60527	Oligonucleotide pr

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C 293	8.4	49.4	10	1	ADU182319	Human neurotrophin
C 294	8.4	49.4	10	1	ADU18500	Hypoxia-related tu
C 295	8.4	49.4	10	1	ADU18776	Hypoxia-related tu
C 296	8.4	49.4	10	1	ADU18606	Hypoxia-related tu
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C 301	8.4	49.4	11	1	ABV64000	Human skin EST 178
C 302	8.4	49.4	11	1	ABV66421	Human skin EST 420
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C 310	8.4	49.4	12	1	AAH80290	Oligo HCV-139, tar
C 311	8.4	49.4	12	1	AAZ41785	Organic material d
C 312	8.4	49.4	12	1	AAZ41569	Microbe detection
C 313	8.4	49.4	12	1	AAZ88808	Gamma-conopeptide
C 314	8.4	49.4	12	1	AAZ34552	Purative Alfini bi
C 315	8.4	49.4	12	1	AAZ34552	Primer used to ill
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1	ABI50359	Oligonucleotide pr
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1	ABI37591	Oligonucleotide pr
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1	ABI23840	Oligonucleotide pr
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1	ABI73498	Oligonucleotide pr
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1	ABI54050	Oligonucleotide pr
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1	ABI62847	Oligonucleotide pr
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1	ABI33500	Oligonucleotide pr
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1	ABI64366	Oligonucleotide pr
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1	ABI80201	Oligonucleotide pr
1	ABI02907	Oligonucleotide pr
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1	ABI42471	Oligonucleotide pr
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 C 448 8.4 49.4 12 1 ABS65874 Chromosomal abnorm
 C 449 8.4 49.4 12 1 ADF78583 Human chromosome 2
 C 450 8.4 49.4 12 1 ADR98159 Human DNA PCR prim
 C 451 8.4 49.4 12 1 ADS08846 Zinc finger protei
 C 452 8 47.1 9 1 ABQ72166 Zinc finger protei
 C 453 8.4 49.4 12 1 ABI75722 Zinc finger target
 C 454 8 47.1 9 1 ADA64492 Zinc finger target
 C 455 8 47.1 9 1 ADA64493 Zinc finger target
 C 456 8 47.1 9 1 ADM23185 Synthetic zinc fin
 C 457 8 47.1 9 1 ADM23184 Synthetic zinc fin
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 C 464 8 47.1 10 1 AAF34623 Yeast NORF gene SA
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ALIGNMENTS

RESULT 1

AAT89947
 ID AAT89947 standard; DNA; 17 BP.

XX AAT89947;

XX 05-MAR-1998 (first entry)

XX Human Interleukin-1 receptor antagonist intron 2 PCR primer 1.

XX Interleukin-1 receptor antagonist; IL-1ra; ulcerative colitis; diagnosis; prognosis; inflammatory bowel disease; PCR primer; ss.

XX Synthetic.

OS Homo sapiens.

XX WO9725445-A1.

XX 17-JUL-1997.

XX 08-JAN-1997; 97WO-US000042.

XX 12-JAN-1996; 96US-00587911.

XX (CEDA-) CEDARS SINAI MEDICAL CENT.

XX (UYVI-) UNIV VIRGINIA PATENT FOUND.

Human AKR1B1 gene
 Human CFL1 primer
 Human prothrombin
 Hypoxia-related tu
 Bovine myostatin m
 Human skin EST 244
 Human skin EST 435
 Human skin EST 960
 Human skin EST 373
 Human skin EST 218
 Human facial skin-
 Human facial skin-
 Human nicking agen
 Human nicking agen
 Human nicking agen
 Human nicking agen
 Human SNP detectio
 Human skin stress/
 Human skin stress/
 Human skin EST 380
 Human skin EST 769
 Human skin EST 819
 Human skin EST 271
 Human skin EST 776
 Human skin EST 234
 Human skin EST 176
 Human skin EST 622
 Human skin EST 847
 Human skin EST 105
 Human skin EST 307
 Human skin EST 977
 Human skin EST 933
 Human skin EST 191
 Human skin EST 627
 Human skin EST 759
 Rat VR1 exon la tr
 Human hair-bearing
 Human hair-bearing
 Human facial skin-
 Human facial skin-
 Human facial skin-

472 8 47.1 10 1 ABL01209
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 C 501 7.8 45.9 11 1 ABV65285
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 C 511 7.8 45.9 11 1 ADQ32561
 C 512 7.8 45.9 11 1 ADQ33791
 C 513 7.8 45.9 11 1 ADQ32382

XX Cominelli F, Pizarro T, Rotter JI, Yang H;
 PI WPI; 1997-372880/34.
 DR Screening for ulcerative colitis in subjects of Jewish ancestry - by
 PT detecting allele 2 of the VNTR (variable number of tandem repeats)
 PT polymorphism at intron 2 of the IL-1 receptor antagonist gene.
 XX Claim 7; Page 17; 22pp; English.
 XX This PCR primer and primer AAT89409 are used to amplify intron 2 of the
 CC human interleukin-1 receptor antagonist gene (IL-1ra) in a novel method
 CC to screen for ulcerative colitis (UC) in a subject of Jewish ancestry.
 CC There is an association between allele 2 of the variable number of tandem
 CC repeats (VNTR) polymorphism at intron 2 of IL-1ra, an important
 CC endogenous regulator of inflammation, and UC in humans of Jewish
 CC ancestry. This method can be used for the diagnosis and prognosis of UC
 CC in Jewish patients for UC and distinguishing UC from Crohn's disease (CD)
 CC and other inflammatory disease of the bowel
 XX Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;
 SQ Query Match 100.0%; Score 17; DB 1; Length 17;
 Best Local Similarity 100.0%; Pred. No. 15;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTCAGCAACACTCTCTAT 17
 Db |||||
 1 CTCAGCAACACTCTCTAT 17
 RESULT 2
 AAT89409
 ID AAT89409 standard; cDNA; 17 BP.
 AC AAT89409;
 XX 22-APR-1998 (first entry)
 DT Human IL-1RN gene intron 2 PCR primer 1.
 DE Osteoporosis; interleukin-1 receptor antagonist; IL-1RN; allele;
 KW bone mineral density; post-menopause; PCR primer; ss.
 XX Synthetic.
 OS Homo sapiens.
 XX WO9738135-A1.
 PN 16-OCT-1997.
 PD 03-APR-1997; 97WO-US005626.
 PF 05-APR-1996; 96US-00628282.
 PR (MEDI-) MEDICAL SCI SYSTEMS INC.
 XX Duff GW, Russell G, Eastell R;
 PI WPI; 1997-512741/47.
 DR Detecting genetic predisposition for osteoporosis - by detecting
 XX interleukin-1 receptor antagonist gene IL-1RN allele 2 in the genomic DNA
 PT of a patient.
 PT Claim 2; Page 9; 21pp; English.
 PS PCR primers AAT89409 and AAT89410 are used to amplify a region of the
 CC interleukin-1 receptor antagonist gene, IL-1RN, intron 2 which contains a
 CC variable number tandem repeat (VNTR) region that gives rise to five
 CC alleles. This product is used for predicting the risk of osteoporosis in
 CC a subject by determining the allelic and genetic polymorphism pattern for

CC IL-1RN in genomic DNA. A pattern of at least one copy of the IL-1RN
 CC allele 2 indicates an increased susceptibility to osteoporosis. The
 CC methods can predict low bone mineral density (BMD) and the rate of bone
 CC density loss and thereby a susceptibility to osteoporosis. Individuals so
 CC identified can then be treated more aggressively to prevent or retard the
 CC occurrence of disease
 XX Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;
 SQ Query Match 100.0%; Score 17; DB 1; Length 17;
 Best Local Similarity 100.0%; Pred. No. 15;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTCAGCAACACTCTCTAT 17
 Db |||||
 1 CTCAGCAACACTCTCTAT 17
 RESULT 3
 AAV62390
 ID AAV62390 standard; DNA; 17 BP.
 XX AAV62390;
 AC AAV62390;
 XX 19-JAN-1999 (first entry)
 DT IL-1 receptor antagonist gene intron 2 PCR primer #1.
 DE Human; interleukin-1 receptor antagonist gene; IL-1; polymorphism;
 KW diagnosis; osteoporosis; PCR primer; ss.
 XX Synthetic.
 OS Homo sapiens.
 XX WO9844150-A1.
 PN 08-OCT-1998.
 PD 27-MAR-1998; 98WO-GB000944.
 PF 27-MAR-1997; 97GB-00006359.
 PR (GEMI-) GEMINI RES LTD.
 PA Keen RW, Spector TD;
 PI WPI; 1998-557135/47.
 DR Diagnosis of osteoporosis by determining genotype of interleukin-1
 PT receptor antagonist gene - useful for diagnosing patient pre-disposition
 PT or susceptibility to osteoporosis and for therapeutic intervention.
 XX Claim 9; Page 10; 36pp; English.
 PS A method has been developed for the diagnosis of osteoporosis comprising
 CC determining the genotype of an interleukin-1 (IL-1) receptor antagonist
 CC gene (IL-1RN). The present sequence represents a PCR primer adapted to
 CC amplify a portion of intron 2 of an IL-1RN for use in the method of the
 CC invention. The method can be used for the diagnosis of disease, including
 CC diagnosis of osteoporosis and predisposition or susceptibility to
 CC osteoporosis and for therapy
 XX Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;
 SQ Query Match 100.0%; Score 17; DB 1; Length 17;
 Best Local Similarity 100.0%; Pred. No. 15;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTCAGCAACACTCTCTAT 17
 Db |||||
 1 CTCAGCAACACTCTCTAT 17

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RESULT 4
AAV60230
ID AAV60230 standard; DNA; 17 BP.
XX AC AAV60230;
XX DT 25-NOV-1998 (first entry)
XX DE PCR primer used to amplify interleukin-1 receptor antagonist (IL-1RN).
XX KW Interleukin-1 receptor antagonist; IL-1RN; predisposition;
XX KW coronary artery disease; screen; PCR primer; ss.
XX YX Synthetic.
XX PN WO9840517-A1.
XX PD 17-SEP-1998.
XX PF 09-MAR-1998; 98WO-US004725.
XX PR 10-MAR-1997; 97US-00813456.
XX PA (MEDI-) MEDICAL SCI SYSTEMS INC.
XX PI Francis SE, Crossman DC, Duff GW;
XX DR WPI; 1998-520829/44.
XX PT Detection of predisposition to coronary artery disease - by comparative
XX PT measurement of levels of expression of alleles from the interleukin 1
XX PT locus.
XX PS Claim 6; Page 15; 22pp; English.
XX CC PCR primers AAV60230-31 were used to amplify alleles associated with the
XX CC interleukin-1 receptor antagonist (IL-1RN). The specification describes a
XX CC method for determination of a patient's predisposition to coronary artery
XX CC disease. The method comprises comparing an allele with a second allele
XX CC which is predictive of coronary artery disease, where similarity between
XX CC the first and second alleles indicates a predisposition to coronary
XX CC artery disease. The method is used to genotype an individual's
XX CC interleukin (IL)-1 loci, the overexpression of which correlates with
XX CC coronary artery disease. The method is used to screen a patients'
XX CC predisposition to coronary artery disease
XX SQ Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;

Query Match 100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 5
AAV32393
ID AAV32393 standard; DNA; 17 BP.
XX AC AAV32393;
XX DT 11-SEP-1998 (first entry)
XX DE Interleukin-1RN primer 1.
XX KW Genetic polymorphism; PCR; primer; amplification; interleukin-1RN;
XX KW sight threatening diabetic retinopathy; interleukin-1-alpha; IL-1RN;
XX KW interleukin-1-beta; ss.
XX YX Synthetic.
XX PN Homo sapiens.
XX PD 20-OCT-1999.

XX PN WO9815653-A1.
XX PD 16-APR-1998.
XX PF 09-OCT-1997; 97WO-GB002790.
XX PR 10-OCT-1996; 96GB-00021129.
XX PA (DUFF/) DUFF G.
XX PA (RENN/) RENNIE I.
XX PA (RICH/) RICHARDSON R.
XX PI Duff G, Rennie I, Richardson R;
XX DR WPI; 1998-240835/21.
XX PT Predicting increased risk of sight-threatening diabetic retinopathy -
XX PT comprises identifying genetic polymorphism pattern for genes IL-1A, IL-1B
XX PT and IL-1RN, useful to allow treatment before clinical symptoms occur.
XX PS Claim 2; Page 33; 41pp; English.
XX CC Interleukin-1RN (IL-1RN) primers 1 and 2 (AAV32394) were used to amplify
XX CC the IL-1RN gene region to identify polymorphism of the VNTR region at the
XX CC IL-1RN intron 2 locus. The invention claims to provide a method for
XX CC predicting the risk of sight threatening diabetic retinopathy. The method
XX CC involves isolating DNA from a patient and determining the DNA
XX CC polymorphism pattern of the genes that code for interleukin-1-alpha,
XX CC interleukin-1-beta and interleukin-1RN. The polymorphic pattern
XX CC identified is then compared with controls of known DNA polymorphism
XX CC patterns thereby identifying patients carrying a genetic polymorphism
XX CC associated with increased risk of sight threatening diabetic retinopathy.
XX CC The method may be able to identify diabetic patients at risk before the
XX CC clinically detectable disorders occur. Polymorphism pattern determination
XX CC of IL genes involved PCR reactions using primers AAV32399-V32398. The
XX CC method is also claimed to be useful in conjunction with identification of
XX CC other genes associated with sight threatening diabetic retinopathy in
XX CC genomic DNA and therefore, in identifying diabetic patients expressing
XX CC multiple risk patterns
XX SQ Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;

Query Match 100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 6
AAZ10706
ID AAZ10706 standard; DNA; 17 BP.
XX AC AAZ10706;
XX DT 23-NOV-1999 (first entry)
XX DE PCR primer 1 used to amplify part of the interleukin-1RN gene.
XX KW Polymorphism; interleukin-1RN; IL-1RN; chronic ulcer;
XX KW inflammatory cytokine; dermal ulcer; venous ulcer; pressure sore;
XX KW decubitis ulcer; PCR primer; ss.
XX YX Synthetic.
XX OS Homo sapiens.
XX PN GB2336431-A.
XX PD 20-OCT-1999.

```

PF 17-APR-1998; 98GB-00008202.
 XX
 PR 17-APR-1998; 98GB-00008202.
 XX
 XX (JOHJ) JOHNSON & JOHNSON MEDICAL LTD.
 XX
 XX Harvey W;
 XX
 XX WPI; 1999-543555/46.
 XX
 XX Polymorphism typing for genes encoding inflammatory cytokines useful for
 PT the analysis of chronic ulcers.
 PT
 XX Example; Page 19; 25pp; English.
 PS
 XX PCR primers AAZ10706-07 were used to identify a polymorphism in intron 2
 CC of the interleukin-1RN (IL-1RN) gene, in the method of the invention. The
 CC specification describes a method for diagnosing the susceptibility to
 CC chronic ulcers, predicting their severity and/or the efficacy of the
 CC healing response generated by the body comprises polymorphism typing for
 CC genes encoding inflammatory cytokines. The method can be used for the
 CC analysis of dermal ulcers e.g. venous ulcers, pressure sores or decubitis
 CC ulcers. The analysis can be used to provide an early diagnosis of the
 CC predisposition to developing a chronic ulcer. Patients can be supervised
 CC over a period of time at which they are thought to be at risk from
 CC developing a chronic ulcer to allow early diagnosis and preventative
 CC intervention before the clinical symptoms are noticed. Individual
 CC treatments can then be used to suit patients with subtle differences in
 CC their disease state
 XX
 SQ Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;
 Query Match 100.0%; Score 17; DB 1; Length 17;
 Best Local Similarity 100.0%; Pred. No. 15;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTCAGCAACACTCCTAT 17
 DB 1 CTCAGCAACACTCCTAT 17
 RESULT 7
 AAX75917
 ID AAX75917 standard; DNA; 17 BP.
 XX
 AC AAX75917;
 XX
 DT 29-JUL-1999 (first entry)
 XX
 DE Human interleukin 1 PCR primer SEQ ID NO:5.
 XX
 KW Human; interleukin 1; IL-1B; IL-1A; IL-1RN; diagnosis; detection;
 KW chronic obstructive airway disease; chronic bronchitis; emphysema;
 KW asthma; chronic bronchiolitis; proinflammatory haplotype; PCR primer; ss.
 XX
 XX Synthetic.
 OS Homo sapiens.
 XX
 PN WO924615-A2.
 XX
 PD 20-MAY-1999.
 XX
 PF 09-NOV-1998; 98WO-US023721.
 XX
 PR 07-NOV-1997; 97GB-00023553.
 PR 12-JAN-1998; 98US-00005923.
 XX
 XX (MEDI-) MEDICAL SCI SYSTEMS INC.
 PA
 XX Duff GW, Giovine M, Barnes PJ, Lim S;
 PI WPI; 1999-327420/27.
 XX
 XX
 PT New method of determining a patient's susceptibility to inflammatory
 disorders - by detecting the presence of an IL-1 (44112332) haplotype,
 XX

PT Genotyping nucleic acid samples for interleukin-1 (IL-1) proinflammatory
 PT haplotype alleles, useful for predicting susceptibility to developing
 PT chronic obstructive airway disease.
 XX
 PS Claim 7; Page 20; 37pp; English.
 XX
 XX The present invention describes genotyping a nucleic acid sample from a
 CC subject to determine at least one allele of an interleukin-1 (IL-1)
 CC proinflammatory haplotype. A method has also been described for
 CC determining a subject's susceptibility to developing chronic obstructive
 CC airway disease (COAD) or for predicting the rapidity or ultimate
 CC progression of a COAD in the subject by: (a) obtaining a nucleic acid
 CC sample from the subject; and (b) detecting at least one allele of an IL-1
 CC proinflammatory haplotype in the sample, where detection of at least one
 CC of these alleles indicates that the patient has an increased
 CC susceptibility to developing COAD. The method is useful for determining
 CC the susceptibility of subjects to developing chronic obstructive airway
 CC disease or for predicting the rapidity or ultimate progression of chronic
 CC obstructive airway disease (COAD). COAD can be asthma, emphysema, chronic
 CC bronchitis or chronic bronchiolitis. The method provides for early
 CC identification of chronic obstructive airway disease (COAD), facilitating
 CC administration of appropriate treatment at the earliest stage, thereby
 CC increasing the probability of a positive outcome. The present sequence
 CC represents a PCR primer used in the method of the invention
 XX
 SQ Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;
 Query Match 100.0%; Score 17; DB 1; Length 17;
 Best Local Similarity 100.0%; Pred. No. 15;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTCAGCAACACTCCTAT 17
 DB 1 CTCAGCAACACTCCTAT 17
 RESULT 8
 AAX16621
 ID AAX16621 standard; DNA; 17 BP.
 XX
 AC AAX16621;
 XX
 DT 29-APR-1999 (first entry)
 XX
 DE Interleukin 1 (44112332) haplotype PCR primer #15.
 XX
 KW Interleukin 1; IL-1; haplotype; inflammatory disorder; alopecia areata;
 KW coronary artery disease; osteoporosis; nephropathy; diabetes mellitus;
 KW Graves disease; systemic lupus erythematosus; lichen sclerosis;
 KW ulcerative colitis; PCR primer; ss.
 XX
 XX Synthetic.
 OS Homo sapiens.
 XX
 PN WO9854359-A1.
 XX
 PD 03-DEC-1998.
 XX
 PF 21-MAY-1998; 98WO-GB001481.
 XX
 PR 29-MAY-1997; 97GB-00011040.
 XX
 XX (DUFF/) DUFF G.
 PA (COXA/) COX A.
 PA (CAMP/) CAMP N J.
 PA (DGIO/) DE GIOVINE F S.
 XX
 XX Duff G, Cox A, Camp NJ, De Giovine FS;
 XX WPI; 1999-080814/07.
 DR
 XX New method of determining a patient's susceptibility to inflammatory
 PT disorders - by detecting the presence of an IL-1 (44112332) haplotype,
 PT

PT useful in designing treatment strategies that modulate the activity of
 XX proteins produced by the IL-1 gene cluster.

PS Claim 3; Page 33; 49pp; English.

XX A method has been developed for determining a patient's susceptibility to
 CC an inflammatory disorder. The method comprises the detection of an
 CC interleukin 1 (IL-1) (44112332) haplotype in a sample obtained from the
 CC patient, where its presence indicates susceptibility to an inflammatory
 CC disorder. AAX16607 to AAX16631 represent PCR primer used in the method
 CC for detecting the IL-1 (44112332) haplotype. The method provides kits for
 CC the early prediction of a patient's susceptibility to inflammatory
 CC disorders, including coronary artery disease, osteoporosis, nephropathy
 CC in diabetes mellitus, alopecia areata, Graves disease, systemic lupus
 CC erythematosus, lichen sclerosis and ulcerative colitis. The detection of
 CC alleles of the haplotype can be applied to particular inflammatory
 CC disorders, comprising diabetic retinopathy, juvenile chronic arthritis,
 CC psoriasis, and insulin dependent diabetes. The identification of a
 CC disease-associated haplotype enables determination of which alleles are
 CC causative, and this information is useful in designing treatment
 CC strategies, including gene therapy and treatment using various agents
 CC that modulate the activity of proteins produced by the IL-1 gene cluster.
 CC Some alleles from the IL-1 gene cluster are associated with particular
 CC inflammatory diseases, and insufficient IL-1 production appears to act
 CC centrally in the pathology of these diseases. Therefore, the use of IL-1
 CC gene clusters is useful in determining genetic susceptibility to
 CC inflammatory diseases, including those with a multifactorial etiology
 CC with a polygenic component

XX Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;

Query Match 100.0%; Score 17; DB 1; Length 17;

Best Local Similarity 100.0%; Pred. No. 15;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCTCTAT 17

Db 1 CTCAGCAACACTCTCTAT 17

RESULT 9

ABX15549

ID ABX15549 standard; DNA; 17 BP.

XX AC ABX15549;

DT 11-APR-2003 (first entry)

XX Human IL-1 genotyping marker VNTR primer #1.

XX Human; ss; PCR; primer; interleukin-1; IL-1; marker VNTR; nephropathy;
 KW inflammatory disease; Systemic Inflammatory Response; SIRS; genotyping;
 KW Alzheimer's disease; arthritis; acute joint inflammation; ophthalmopathy;
 KW juvenile chronic arthritis; asthma; bronchial asthma; pulmonary disease;
 KW chronic obstructive airways disease; cardiovascular disease; thyroiditis;
 KW atherosclerosis; autoimmune carditis; cardiomyopathy; ulcerative colitis;
 KW cardiac cell dysfunction; aortic smooth muscle cell activation; trauma;
 KW inflammatory bowel disease; HIV infection; coronary artery lesion;
 KW Kawasaki's syndrome; cervical lymphadenopathy; diabetic nephropathy;
 KW glomerulonephritis; diabetic retinopathy; Grave's ophthalmopathy;
 KW osteoporosis; bone loss; otitis media; pancreatitis; periodontal disease;
 KW chronic lung disease; chronic sinusitis; chronic lymphocytic thyroiditis;
 KW urinary tract infection; chronic prostatitis; immunological disorder;
 KW chronic pelvic pain syndrome; alopecia areata; Grave's disease;
 KW thyroid disease; goiter; struma lymphomatosa; sleep disorder; neoplasia;
 KW chronic fatigue syndrome; obesity; infectious disease; leishmaniasis;
 KW leprosy; myocardial dysfunction; breast cancer; organ transplant;
 KW Hodgkin's disease; hormonal regulation; fertility; septicaemia.

XX Homo sapiens.

XX US2002146700-A1.

XX 10-OCT-2002.

XX 27-APR-2001; 2001US-00845129.

XX 29-MAY-1997; 97GB-00011040.

XX 30-JUN-1999; 99US-00345217.

XX (INTE-) INTERLEUKIN GENETICS INC.

XX Duff GW, Cox A, Camp NJ, Di Giovine FS;

XX WPI; 1999-080814/07.

XX New method of determining a patient's susceptibility to inflammatory
 PT disorders - by detecting the presence of an IL-1 (44112332) haplotype,
 PT useful in designing treatment strategies that modulate the activity of
 PT proteins produced by the IL-1 gene cluster.

XX Claim 5; Page 20; 42pp; English.

XX The invention relates to a method for determining whether a subject has
 CC or is predisposed to developing a disease or condition that is associated
 CC with an IL-1-inflammatory haplotype. The method involves detecting at
 CC least one allele of the haplotype, where the presence of the allele
 CC indicates that the subject is predisposed to the development or has the
 CC disease or condition. The invention allows the determination of an
 CC individual's likelihood for developing a particular disease or condition
 CC associated with interleukin 1 (IL-1) polymorphisms without necessarily
 CC determining or characterising the causative genetic variation. Diseases
 CC such as inflammatory disease e.g. Systemic Inflammatory Response (SIRS),
 CC Alzheimer's disease; arthritis e.g. acute joint inflammation, juvenile
 CC chronic arthritis; asthma e.g. bronchial asthma, chronic obstructive
 CC airways disease; cardiovascular diseases e.g. atherosclerosis, autoimmune
 CC carditis; cardiomyopathy and cardiac cell dysfunction e.g. aortic smooth
 CC muscle cell activation, cardiac cell apoptosis; gastrointestinal
 CC inflammations e.g. inflammatory bowel disease, ulcerative colitis; HIV
 CC infection; Kawasaki's syndrome e.g. cervical lymphadenopathy, coronary
 CC artery lesions; nephropathies e.g. diabetic nephropathy,
 CC glomerulonephritis; ophthalmopathies e.g. diabetic retinopathy, Grave's
 CC ophthalmopathy; osteoporosis e.g. bone loss, otitis media; pancreatitis;
 CC periodontal disease; pulmonary diseases e.g. chronic lung disease,
 CC chronic sinusitis; thyroiditis e.g. chronic lymphocytic thyroiditis;
 CC urinary tract infections e.g. chronic prostatitis, chronic pelvic pain
 CC syndrome; immunological disorders e.g. alopecia areata, Graves disease;
 CC thyroid diseases e.g. goiter, struma lymphomatosa; sleep disorders;
 CC chronic fatigue syndrome; obesity; infectious diseases e.g. leprosy,
 CC leishmaniasis; trauma e.g. cerebral trauma, myocardial dysfunction;
 CC neoplasias e.g. breast cancer, Hodgkin's disease; hormonal regulation e.g.
 CC fertility, septicaemia; organ transplants. This allows for a more
 CC customised approach to preventing the onset or progression of the disease
 CC or condition, e.g. a clinician can more effectively prescribe a therapy
 CC that will address the molecular basis of the disease or condition. The
 CC present sequence represents the sequence of the human IL-1 genotyping
 CC marker VNTR primer #1

XX Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;

Query Match 100.0%; Score 17; DB 1; Length 17;

Best Local Similarity 100.0%; Pred. No. 15;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCTCTAT 17

Db 1 CTCAGCAACACTCTCTAT 17

RESULT 10

AAZ37216

ID AAZ37216 standard; DNA; 17 BP.

XX AC AAZ37216;

DT 28-JAN-2000 (first entry)

XX PCR primer for interleukin-1RN (VNTR).

DE PCR primer; IL-1; IL-1A; IL-1B; TNFA; interleukin-1; foetus;

XX PCR primer; IL-1; IL-1A; IL-1B; TNFA; interleukin-1; foetus;

KW adverse pregnancy outcome; allele detection; low birth weight; LBW;

KW pre-term baby; premature baby; tumour necrosis factor-alpha; ss.

XX Synthetic.

OS Homo sapiens.

XX WO9954707-A2.

PN 28-OCT-1999.

PD 21-APR-1999; 99WO-US008794.

XX 21-APR-1999; 98US-0082487P.

XX (MEDI-) MEDICAL SCI SYSTEMS INC.

PA Kornman KS, Offenbacher S, Duff GW;

XX WPI; 2000-013279/01.

DR Fetal testing for prediction of low birth weight by detecting an

XX interleukin allele.

PT Claim 5; Page 73; 78pp; English.

XX This sequence represents a PCR primer for interleukin-1RN (IL-1RN). The

CC invention relates to a method for determining whether a foetus is

CC predisposed to having an adverse pregnancy outcome by detecting an

CC interleukin-1A (IL-1A) (+4845) allele 2, an IL-1 (-511) allele 1 or an

CC allele in linkage disequilibrium with either of these alleles. The method

CC comprises: (a) obtaining a nucleic acid sample from the subject; and (b)

CC detecting an IL-1A (+4845) allele 2, an IL-1 (-511) allele 1 or an allele

CC in linkage disequilibrium with either of these alleles, where detection

CC of the alleles or an allele in linkage disequilibrium with them indicates

CC that the fetus is predisposed to an adverse pregnancy outcome. The method

CC is used to determine whether a fetus is predisposed to having a low birth

CC weight (LBW). By determining the IL-1 or tumour necrosis factor-alpha

CC (TNF-A) genotype an appropriate therapeutic, such as a corticosteroid,

CC antimetabolite, cytotoxic drug, colchicine or anti-cytokine, can be

CC administered to compensate for the LBW causative functional mutation. An

CC appropriate therapeutic can be used to treat a subject predisposed to

CC having a LBW baby, especially a pre-term or premature baby

XX Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;

SQ Query Match 100.0%; Score 17; DB 1; Length 17;

Best Local Similarity 100.0%; Pred. No. 15;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGCAACACTCCTAT 17

Db 1 CTCAGCAACACTCCTAT 17

RESULT 11

AAA61915

ID AAA61915 standard; DNA; 17 BP.

XX AAA61915;

AC 20-NOV-2000 (first entry)

XX Human IL-1RN (VNTR) polymorphic locus PCR primer, SEQ ID NO:1.

DT Human; IL-1 gene cluster; interleukin-1; chromosome 2q13-14;

XX Human; IL-1 gene cluster; interleukin-1; chromosome 2q13-14;

DE proinflammatory; cytokine; systemic inflammatory response; sepsis;

KW dysregulation; pattern 2 allele; genotyping; IL-1RN gene;

KW IL-1 receptor antagonist; IL-1RN (VNTR) polymorphic locus;

KW variable number of tandem repeats; PCR primer; ss.

XX Homo sapiens.

XX WO200037679-A2.

XX 29-JUN-2000.

PD 01-NOV-1999; 99WO-US025633.

XX 30-OCT-1998; 98US-00183850.

XX (INTE-) INTERLEUKIN GENETICS INC.

PA Di Giovine FS, Duff GW;

PI WPI; 2000-442691/38.

XX Determining susceptibility to developing sepsis for preventing or

XX treating sepsis, involves detecting an allele of interleukin-1 genetic

PT pattern that leads to dysregulated inflammatory response.

XX Disclosure; Page 35; 66pp; English.

XX The invention relates to methods of determining an individual's

CC susceptibility to developing and/or rapidly progressing into sepsis via

CC genotyping of a member of the IL-1 (interleukin-1) gene cluster. The

CC method comprises obtaining an IL-1 gene cluster nucleic acid from an

CC individual and determining whether that individual's DNA contains at

CC least one allele of an IL-1 genetic pattern that leads to a dysregulated

CC inflammatory response. The presence of a pattern 2 allele or a marker in

CC linkage disequilibrium with a pattern 2 allele in the sample, indicates

CC that the patient has an increased susceptibility to developing sepsis.

CC The IL-1 gene cluster comprises a region of 430 kb which is located on

CC the long arm of chromosome 2 (2q13-14). It contains the IL-1A gene which

CC encodes IL-1-alpha, the IL-1B gene which encodes IL-1-beta, and the IL-

CC 1RN gene which encodes the IL-1 receptor antagonist. IL-1 is a

CC proinflammatory cytokine released by macrophages which can induce a

CC systemic response to local injury or infection. It thus plays a key role

CC in the development of sepsis, which can lead to intravascular

CC coagulation, multiple organ failure, cardiovascular collapse, and death.

CC It is therefore useful to identify those individuals who are at risk of

CC developing an exaggerated systemic inflammatory response to septic

CC stimuli, so that sepsis therapy (e.g. with IL-1 modulators) can be

CC tailored to the individual's genetic profile. Sequences AAA61915-A61928

CC represent PCR primers which may be used to amplify IL-1 gene cluster

CC polymorphic loci. Sequences AAA61915-A61916 are primers used to amplify

CC and type the IL-1RN (VNTR) (variable number of tandem repeats)

CC polymorphic locus

XX Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;

SQ Query Match 100.0%; Score 17; DB 1; Length 17;

Best Local Similarity 100.0%; Pred. No. 15;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGCAACACTCCTAT 17

Db 1 CTCAGCAACACTCCTAT 17

RESULT 12

AA63787

ID AA63787 standard; DNA; 17 BP.

XX AA63787;

AC 08-FEB-2001 (first entry)

XX Human IL-1RN gene VNTR primer #1.

DE Human; IL-1RN; interleukin-1 receptor; cytostatic; antiinflammatory;

KW immunosuppressive; dermatological; antimicrobial; antiarthritic;

KW

KW IL-1 receptor antagonist;
 KW tumour necrosis factor alpha antagonist interstitial lung disease;
 KW interstitial pneumonia; pulmonary fibrosis; rheumatoid arthritis;
 KW systemic lupus erythematosus; Sjogren's syndrome; systemic sclerosis;
 KW dermatomyositis; chromosome 2; primer; ss.
 XX Homo sapiens.
 XX WO200060117-A2.
 XX 12-OCT-2000.
 XX 31-MAR-2000; 2000WO-US008492.
 XX 02-APR-1999; 99US-00286108.
 XX (INTE-) INTERLEUKIN GENETICS INC.
 XX Duff GW, Di Giovine FS, Whyte M;
 XX WPI; 2000-656234/63.
 XX Method for predicting the risk of interstitial lung disease, comprising
 PT detecting an interleukin-1 receptor antagonist allele and tumor necrosis
 PT alpha allele or an allele in linkage disequilibrium with either of these
 PT alleles.
 XX
 XX Example 2; Page 72; 102pp; English.
 XX
 CC The present sequence is provided in a specification relating to a method
 CC for determining whether a subject has or is predisposed to develop an
 CC interstitial lung disease. The method involves detecting an interleukin-1
 CC receptor antagonist (IL-1RN) (+2018) allele 2, a tumour necrosis alpha
 CC (TNF-A)(-308) allele 2, or an allele in linkage disequilibrium with
 CC either of these two alleles. The method may be used to determine whether
 CC a subject has or is predisposed to develop an interstitial pneumonia or a
 CC pulmonary fibrosis and other disorders such as rheumatoid arthritis,
 CC systemic lupus erythematosus, Sjogren's syndrome, systemic sclerosis,
 CC dermatomyositis. The method is also used for identifying molecules which
 CC can be used as therapeutics for treating interstitial lung disease
 XX
 XX Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;

Query Match 100.0%; Score 17; DB 1; Length 17;
 Best Local Similarity 100.0%; Pred. No. 15;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
 |||||
 Db 1 CTCAGCAACACTCCTAT 17

RESULT 13
 AAF27686
 ID AAF27686 standard; DNA; 17 BP.
 XX
 AC AAF27686;
 XX
 DT 02-APR-2001 (first entry)
 XX
 DE Primer #19.
 XX
 XX IL-1; interleukin; inflammation; infection; ss.
 XX Unidentified.
 XX
 XX WO200100880-A2.
 XX
 XX 04-JAN-2001.
 XX
 XX 30-JUN-2000; 2000WO-US018318.
 XX
 XX 30-JUN-1999; 99US-00345217.

XX (INTE-) INTERLEUKIN GENETICS INC.
 XX Duff GW, Cox A, Camp NJ, Di Giovine FS;
 XX WPI; 2001-102903/11.

XX Determining whether a subject has or is predisposed to disease associated
 PT with IL-1 polymorphism involves determining presence of marker or allele
 PT comprising IL-1 inflammatory haplotype.

XX Claim 5; Page 48; 84pp; English.

XX The present invention relates to a new method for determining whether a
 CC subject has or is predisposed to developing a disease or condition that
 CC is associated with an IL (interleukin)-1 inflammatory haplotype,
 CC comprises detecting at least one allele of the haplotype, where the
 CC presence of the allele indicates that the subject is predisposed to the
 CC development or has the disease or condition. The method is useful for
 CC determining whether a subject has or is predisposed to inflammatory
 CC disease, a degenerative disease, an immunological disorder, an infectious
 CC disease, trauma induced disease, or cancer. The above conditions
 CC associated with an IL-1 inflammatory haplotype can be treated or
 CC prevented by administering a therapeutic that compensates for a causative
 CC mutation that is in linkage disequilibrium with at least one IL-1
 CC polymorphism

XX Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;

Query Match 100.0%; Score 17; DB 1; Length 17;
 Best Local Similarity 100.0%; Pred. No. 15;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
 |||||
 Db 1 CTCAGCAACACTCCTAT 17

RESULT 14
 AAC89171
 ID AAC89171 standard; DNA; 17 BP.

XX AAC89171;

XX 08-MAR-2001 (first entry)

XX Human IL-1RN (VNTR) polymorphic locus PCR primer #1.

XX Human; PCR primer; cardiant; vasotropic; polymorphic; chromosome 2q13;
 KW interleukin-1 gene cluster; IL-1; cardiovascular disorder; IL-1RN;
 KW fragile plaque disorder; occlusive disorder; in-stent restenosis; ss.

XX Homo sapiens.

XX WO200072015-A2.

XX 30-NOV-2000.

XX 26-MAY-2000; 2000WO-US014775.

XX 26-MAY-1999; 99US-00320395.

XX 01-NOV-1999; 99US-00431352.

XX (INTE-) INTERLEUKIN GENETICS INC.

XX Francis SE, Crossman DC, Duff GW, Kornman KS;

XX WPI; 2001-032066/04.

XX Diagnosing cardiovascular disease, or susceptibility, useful e.g. for
 PT selecting treatment, from presence of disease-associated allele,
 PT particularly in interleukin-1 genes.

PS Claim 11; Page 79; 121pp; English.

XX There is an association of patterns of alleles at four polymorphic loci

CC in the interleukin-1 (IL-1) gene cluster with cardiovascular disorders.

CC The present invention relates to a method for the diagnosis of a

CC cardiovascular disorder (CVD), via detecting a CVD-associated allele in a

CC patient sample. The method is used to select an appropriate therapy for

CC CVD, particularly fragile plaque disorder, occlusive disorders or in-

CC stent restenosis, or a condition associated with a mutation with linkage

CC disequilibrium with a CVD-associated allele, or to identify risk factors

CC for CVD, and to formulate a treatment designed to reduce the risk. The

CC present sequence is a PCR primer used in the present invention. The IL-1

CC gene cluster is found at the 2q13 region of chromosome 2

XX

XX Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;

Query Match 100.0%; Score 17; DB 1; Length 17;

Best Local Similarity 100.0%; Pred. No. 15;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCTCTAT 17

Db 1 CTCAGCAACACTCTCTAT 17

RESULT 15

AAAC91425

ID AAD27377 standard; DNA; 17 BP.

XX

XX AAC91425;

XX

XX 20-MAR-2001 (first entry)

XX

XX Human IL-1RN (VNTR) polymorphic locus 5' PCR primer.

DE Human; IL-1A; interleukin-1alpha; IL-1B; interleukin-1beta; IL-1RN;

KW interleukin-1 receptor antagonist; vasotropic; antiinflammatory;

KW hypotensive; anticoagulant; antilipemic; arterial restenosis;

KW restenosis associated allele; RAA; occlusive cardiovascular disorder;

KW restenosis detection; PCR primer; ss.

XX

OS Homo sapiens.

XX

XX WO200071753-A2.

XX

XX 30-NOV-2000.

XX

XX 24-MAY-2000; 2000WO-US014299.

XX

XX 24-MAY-1999; 99US-00317674.

PR 01-NOV-1999; 99US-00431352.

XX

XX (INTE-) INTERLEUKIN GENETICS INC.

XX

XX Korman KS, Duff GW, Crossman DC, Francis SE, Stephenson K;

PI WPI; 2001-025173/03.

XX

XX Diagnosing or determining susceptibility to developing restenosis

PT involves detecting restenosis associated allele in a nucleic acid sample.

XX

XX Claim 5; Page 11; 129pp; English.

XX

XX The present sequence may be used in a method for determining whether a

CC subject has or is predisposed to developing an arterial restenosis. The

CC method comprises detecting a restenosis associated allele (RAA) in a

CC nucleic acid sample from the subject, where detection of the RAA

CC indicates that the subject has or is predisposed to the development of a

CC restenosis. The restenosis associated allelic pattern permits the

CC diagnosis of occlusive cardiovascular disorder. The diagnosis allows the

CC most suitable treatment methods for restenosis to be used e.g. selecting

CC therapies for initial vascular stenosis most likely to avoid subsequent

CC stenoses. The detection methods identify restenosis therapeutics,

CC

CC agonists and antagonists, (proteins, peptides, peptidomimetics, small

CC molecules or nucleic acids, e.g. anti-sense, ribozyme and triplex nucleic

CC acids) which are used to treat restenosis

XX

XX Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;

Query Match 100.0%; Score 17; DB 1; Length 17;

Best Local Similarity 100.0%; Pred. No. 15;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCTCTAT 17

Db 1 CTCAGCAACACTCTCTAT 17

RESULT 16

AAAD27377

ID AAD27377 standard; DNA; 17 BP.

XX

XX AAD27377;

XX

XX 18-APR-2002 (first entry)

XX

XX PCR primer #1, used for genotyping human IL-1RN (VNTR) marker.

DE Human; interleukin-1; inflammatory disorder; coronary artery disease;

KW periodontal disease; Alzheimer's disease; atherosclerosis; osteoporosis;

KW immune response; insulin-dependent diabetes; diabetic retinopathy;

KW renal disease; diabetic nephropathy; hepatic fibrosis; alopecia areata;

KW Graves disease; Graves ophthalmopathy; systemic lupus erythematosus;

KW extrathyroid disease; lichen sclerosis; ulcerative colitis; asthma;

KW rheumatoid arthritis; gastric cancer; juvenile chronic arthritis;

KW interstitial lung disease; idiopathic pulmonary fibrosis; sepsis;

KW multiple sclerosis; acne; IL-1 receptor antagonist; IL-1RN; VNTR;

KW cardiant; dermatological; nootropic; neuroprotective; osteopathic;

KW ophthalmological; PCR primer; ss.

XX

OS Homo sapiens.

XX

XX WO200200933-A2.

XX

XX 03-JAN-2002.

XX

XX 22-JUN-2001; 2001WO-US020079.

XX

XX 23-JUN-2000; 2000US-0213853P.

PR

XX (INTE-) INTERLEUKIN GENETICS INC.

XX

XX Duff GW, Kornman KS;

PI WPI; 2002-139934/18.

XX

XX Screening a substance in a subject for modulating an immune response,

PT comprises genotyping to identify the test subject, and observing a

PT biomarker before and after contacting the subject with the test

PT substance.

XX

XX Example; Page 40; 54pp; English.

XX

XX The present invention relates to methods for identifying a test substance

CC that modulate the immune response in a genotype specific manner. Methods

CC of the invention involve genotyping subjects to identify those having a

CC genotype (e.g. interleukin-1; IL-1) associated with one or more

CC inflammatory disorder. The method comprises genotyping a subject having

CC an inflammatory disease-associated genotype and observing a biomarker in

CC the subject before and after the subject is contacted with the test

CC substance. The methods or cells associated with inflammatory diseases are

CC useful for identifying a substance that is likely to prevent or diminish

CC a specific biological response in subjects having inflammatory disease-

CC associated genotype, where the genotype is associated a pre-disposition

CC to one or more of periodontal disease, coronary artery disease,

CC Alzheimer's disease, atherosclerosis, osteoporosis, insulin- dependent

CC diabetes, diabetic retinopathy, end-stage renal disease, diabetic
 CC nephropathy, hepatic fibrosis, alopecia areata, Graves disease, Graves
 CC ophthalmopathy, extrathyroid disease, systemic lupus erythematosus,
 CC lichen sclerosus, rheumatoid arthritis, juvenile chronic arthritis,
 CC gastric cancer, ulcerative colitis, asthma, interstitial lung disease,
 CC multiple sclerosis, idiopathic pulmonary fibrosis, sepsis and acne. The
 CC invention also relates to a kit comprising primers for the identification
 CC of one or more IL-1 polymorphisms. The present sequence is a PCR primer
 CC which is used for amplifying IL-1 receptor antagonist (IL-1RN; VNPR)
 CC gene. This primer is used in the exemplification of the invention for
 CC genotyping IL-1RN marker
 XX
 SQ Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;
 Query Match 100.0%; Score 17; DB 1; Length 17;
 Best Local Similarity 100.0%; Pred. No. 15;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTCAGCAACACTCCTAT 17
 DB 1 CTCAGCAACACTCCTAT 17
 RESULT 17
 AAL54501
 ID AAL54501 standard; DNA; 17 BP.
 XX
 AC AAL54501;
 DT 16-APR-2003 (first entry)
 XX
 DE Ageing-related condition detection PCR primer, SEQ ID No 7.
 XX
 DE Early onset; progression; ageing-related condition; BOA; allele;
 KW interleukin; (IL)-1; pattern 1; pattern 2; pattern 3; osteoporosis;
 KW osteoarthritis; wrinkled skin; age-related cancer; lifestyle; exercise;
 KW diet; nutraceutical; PCR; primer; ss.
 XX
 OS Unidentified.
 XX
 XX WO2002103031-A2.
 XX
 XX 27-DEC-2002.
 XX
 XX 17-JUN-2002; 2002WO-US019205.
 XX
 XX 15-JUN-2001; 2001US-0298493P.
 XX
 XX (INTE-) INTERLEUKIN GENETICS INC.
 XX
 PI Barnett K, Crossman DC, Duff GW, Francis SE, Kornman KS;
 XX
 XX WPI; 2003-167530/16.
 XX
 XX Determining a subject's susceptibility to an early onset or progression
 XX of an aging-related condition, useful for customizing therapy, comprises
 XX detecting the presence of an allele of an interleukin-1 pattern 1,
 XX pattern 2 and/or pattern 3.
 XX
 XX Example 3; Page 52; 98pp; English.
 XX
 XX The invention relates to a novel method for determining a subject's
 XX susceptibility to the early onset or progression of an ageing-related
 XX condition (EOA). The novel method comprises assessing the subject's
 XX genotype with respect to at least one allele of an interleukin (IL)-1
 XX pattern 1, pattern 2 and/or pattern 3 (the presence or absence of at
 XX least 1 allele provides information about the subject's susceptibility to
 XX an early onset or progression of an ageing-related condition). The method
 XX is useful for determining or predicting a subject's susceptibility to the
 XX early onset or progression of an ageing-related condition (e.g.
 XX osteoporosis, osteoarthritis, wrinkled skin, or age-related cancer) and
 XX for determining an ageing-related phenotype. The method may be a
 XX customised therapy based on the individual's genetic profile, to tailor a

CC recommended lifestyle, including changes in exercise and diet, and to
 CC recommend nutraceuticals that are predicted to benefit a subject having a
 CC particular IL-1 genotype and BOA predisposition. This polynucleotide
 CC sequence represents a PCR primer used in the detection method for
 CC determining EOA predisposition of the invention
 XX
 SQ Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;
 Query Match 100.0%; Score 17; DB 1; Length 17;
 Best Local Similarity 100.0%; Pred. No. 15;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTCAGCAACACTCCTAT 17
 DB 1 CTCAGCAACACTCCTAT 17
 RESULT 18
 AAD51455
 ID AAD51455 standard; DNA; 17 BP.
 XX
 AC AAD51455;
 XX
 DT 16-APR-2003 (first entry)
 XX
 DE Human interleukin-1 (IL-1) gene amplifying PCR primer #7.
 XX
 DE Drug screening; fungicide; gene therapy; antibacterial; infection;
 KW virucide; human; interleukin-1; IL-1; PCR; primer; ss.
 XX
 XX Homo sapiens.
 XX
 XX WO2002101015-A2.
 XX
 XX 19-DEC-2002.
 XX
 XX 11-JUN-2002; 2002WO-US018346.
 XX
 XX 11-JUN-2001; 2001US-0297305P.
 XX
 XX (INTE-) INTERLEUKIN GENETICS INC.
 XX
 PI Dower S, Duff GW;
 XX
 XX WPI; 2003-148793/14.
 XX
 XX New detection reagent, useful for monitoring molecular assembly events to
 XX permit the dissection of genetic and non-genetic influences on biological
 XX activity, comprises an interactive sensor pair.
 XX
 XX Disclosure; Page 41; 56pp; English.
 XX
 XX The invention relates to methods, compositions and apparatus for
 XX monitoring molecular assembly events. It also relates to a detection
 XX reagent comprising an interactive sensor pair. The detection reagent is
 XX useful for monitoring molecular assembly events to permit the dissection
 XX of genetic and non-genetic influences on a particular biological
 XX activity. The method is useful for linking genetic variations to
 XX molecular and physiological events, drug screening, diagnostics, therapy
 XX selection and dosing, patient monitoring or environmental safety. The
 XX interactive sensor pairs may be used to screen for and identify novel
 XX agonists and antagonists or other molecules that modulate a biological
 XX activity. The method is also useful for selecting an appropriate targeted
 XX therapeutic for a subject having an infection, including viral, bacterial
 XX or fungal infection. It is also used in gene therapy. The present
 XX sequence is a PCR primer used for amplifying human interleukin-1 (IL-1)
 XX gene. This sequence is used to illustrate the method of the invention
 XX
 SQ Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;
 Query Match 100.0%; Score 17; DB 1; Length 17;
 Best Local Similarity 100.0%; Pred. No. 15;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGCAACACTCCTAT 17
 DB 1 CTCAGCAACACTCCTAT 17

RESULT 19
 ADL83329
 ID ADL83329 standard; DNA; 17 BP.
 XX AC
 XX ADL83329;
 XX 01-JUL-2004 (first entry)
 XX PCR primer #1 for human IL-1RN (VNTR) DNA.
 XX Single vessel coronary artery disease; SVD; restenosis;
 KW interleukin-1 receptor antagonist; IL-1RN; VNTR; allele; human; PCR;
 KW primer; ss.
 XX Homo sapiens.
 OS US6720141-B1.
 XX 13-APR-2004.
 XX 24-MAY-2000; 2000US-00578534.
 XX 01-NOV-1999; 99US-00431352.
 XX (INTE-) INTERLEUKIN GENETICS INC.
 XX Crossman DC, Duff GW, Francis SE, Kornman KS, Stephenson K;
 PI WPI; 2004-313649/29.
 XX Determining whether a single vessel coronary artery disease (SVD) subject
 PT has or is predisposed to developing restenosis comprises detecting IL-1RN
 PT (VNTR) allele 1 or 2 in a nucleic acid sample from the subject.
 XX Claim 4; SEQ ID NO 7; 76pp; English.

The present invention relates to a method for determining whether a
 CC single vessel coronary artery disease (SVD) subject has or is predisposed
 CC to developing restenosis. The method comprises detecting interleukin-1
 CC receptor antagonist (IL-1RN) (VNTR) allele 1 or 2 in a nucleic acid
 CC sample from the subject, where detection of IL-1RN (VNTR) allele 1 or 2
 CC indicates that the subject has or is predisposed to the development of
 CC restenosis. Also disclosed is a kit for performing the method above. In
 CC the method, the detecting step is selected from allele specific
 CC oligonucleotide (ASO) hybridisation, size analysis, sequencing,
 CC hybridisation, 5' nuclease digestion, single-stranded conformational
 CC polymorphism (SSCP), allele specific hybridisation, primer specific
 CC extension or oligonucleotide ligation assay. The method is useful for
 CC determining whether a single vessel coronary artery disease (SVD) subject
 CC has or is predisposed to developing restenosis. The present sequence
 CC represents a PCR primer used in the method of the invention.

XX Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;
 SQ Query Match 100.0%; Score 17; DB 1; Length 17;
 Best Local Similarity 100.0%; Pred. No. 15;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGCAACACTCCTAT 17
 DB 1 CTCAGCAACACTCCTAT 17

RESULT 20
 ADN48846
 ID ADN48846 standard; DNA; 17 BP.
 XX

AC ADN48846;
 XX 15-JUL-2004 (first entry)
 XX Human secreted interleukin-1RN (IL-1RN) amplifying PCR primer #1.
 DE Early-onset menopause; EOM; diagnosis; therapy; human; interleukin-1RN;
 KW IL-1RN; PCR; primer; ss.
 XX Homo sapiens.
 OS US6730476-B1.
 XX 04-MAY-2004.
 XX 04-AUG-2000; 2000US-00632657.
 XX 30-JUN-1999; 99US-00345217.
 XX (INTE-) INTERLEUKIN GENETICS INC.
 XX Duff G, Kornman K, Van Dijk S;
 PI WPI; 2004-354679/33.
 XX Determining the predisposition to early-onset menopause comprises
 PT detecting in the subject interleukin (IL)-1RN (+2018) allele 2.
 XX Example; SEQ ID NO 9; 57pp; English.
 XX The present invention provides a method for determining the
 CC predisposition of a subject to early-onset menopause (EOM). The invention
 CC is useful in diagnosing, treating and preventing early-onset menopause.
 CC The present sequence is human secreted interleukin-1RN (IL-1RN)
 CC amplifying PCR primer. The sequence is used in exemplification of the
 CC invention.

XX Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;
 SQ Query Match 100.0%; Score 17; DB 1; Length 17;
 Best Local Similarity 100.0%; Pred. No. 15;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGCAACACTCCTAT 17
 DB 1 CTCAGCAACACTCCTAT 17

RESULT 21
 ADO35276
 ID ADO35276 standard; DNA; 17 BP.
 XX ADO35276;
 XX 26-AUG-2004 (first entry)
 XX Human interleukin-1RN (IL-1RN) gene, PCR primer #1.
 DE human; obstructive airway disease; OAD; interleukin-13; IL-1B;
 KW IL-1A; IL-1RN; asthma; interleukin-1A; interleukin-1B; interleukin-1RN;
 KW ss; primer.
 XX Homo sapiens.
 OS US6746839-B1.
 XX 08-JUN-2004.
 XX 01-JUN-2000; 2000US-00584950.
 XX 12-JAN-1998; 98US-00005923.
 XX (INTE-) INTERLEUKIN GENETICS INC.

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XX PI Duff GW, Di Giovine FS, Barnes PJ, Lim S;
XX WPI; 2004-429802/40.
XX
XX PT Determining whether a subject has or is predisposed to developing an
XX obstructive airway disease (OAD), such as asthma, detecting an OAD-
XX associated allele in a nucleic acid sample from the subject, which
XX indicates predisposition to OAD.
XX
XX PS Example 3; SEQ ID NO 13; 61pp; English.
XX
XX CC The invention relates to a method of determining whether a subject has,
XX or is predisposed to developing, an obstructive airway disease (OAD),
XX comprising detecting an OAD-associated allele in a nucleic acid sample
XX from the subject, where the presence of OAD-associated allele indicates
XX that the subject has or is predisposed to the development of OAD. The OAD
XX associated allele is selected from allele 2 of interleukin-13 (IL-13)
XX (-2581), allele 2 of IL-1B (-511), allele 2 of IL-1B (+3954), or an
XX allele in linkage disequilibrium with allele 2 of IL-1B (-511) or allele
XX 2 of IL-1B (+3954) comprising allele 4 of IL-1A (222/223), allele 4 of IL
XX -1A (g25/g26), allele 1 of IL-1A (-889), allele 1 of (+3954), allele 3 of
XX the gaat.p3330 marker, allele 3 of the Y31 marker, allele 2 of IL-1RN
XX (-2018), allele 2 of IL-1RN (VNTR), allele 3 of IL-1A (222/223), allele 3
XX of IL-1A (g25/g26), allele 2 of IL-1A (-889), allele 1 of IL-1B (-511),
XX allele 4 of the gaat.p3330 marker, allele 6 of the Y31 marker, allele 1
XX of IL-1RN (+2018), and allele 1 of IL-1RN (VNTR). The method is useful
XX for determining whether a subject has or is predisposed to developing an
XX OAD, particularly asthma. The method is especially useful for the early
XX identification of those who are generally susceptible to OAD, and those
XX who are susceptible to acute episodes. Early identification would
XX facilitate the prevention or administration of appropriate treatment at
XX the earliest stage, thus increasing the probability of a positive
XX outcome. The present sequence represents a PCR primer used to detect
XX polymorphisms in the human interleukin-1 (IL-1) gene locus.
XX
XX SQ Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;
XX
XX Query Match 100.0%; Score 17; DB 1; Length 17;
XX Best Local Similarity 100.0%; Pred. No. 15;
XX Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
Qy 1 CTCAGCAACACTCTCTAT 17
Db 1 CTCAGCAACACTCTCTAT 17
XXXXXXXXXXXXXXXXXXXX
XXXXXXXXXXXXXXXXXXXX

RESULT 22
ADU16279
ID ADU16279 standard; DNA; 17 BP.
XX
XX AC ADU16279;
XX
XX DT 27-JAN-2005 (first entry)
XX
XX DE Human interleukin-1R genotyping primer seqid 29.
XX
XX KW seronegative spondyloarthropathy; interleukin 1; IL-1;
XX ankylosing spondylitis; interleukin-1R; IL-1R; PCR; primer; ss.
XX
XX OS Homo sapiens.
XX
XX PN WO2004097045-A1.
XX
XX PD 11-NOV-2004.
XX
XX PF 04-MAY-2004; 2004WO-GB001905.
XX
XX PR 01-MAY-2003; 2003GB-00010060.
XX
XX PA (ISIS-) ISIS INNOVATION LTD.
XX
XX PI Brown MA;

XX PI Duff GW, Duff GW, Francis SE, Kornman KS, Stephenson K;
XX WPI; 2004-813241/80.
XX
XX PT New kit comprises a first primer oligonucleotide that hybridizes 5' or 3'
XX to an allele 1 of IL-1A (+4845), IL-1B (-511), IL-1B (+3954), IL-1RN
XX (VNTR), or IL-1RN (+2018), useful for diagnosing, treating, or preventing
XX restenosis.
XX
XX Query Match 100.0%; Score 17; DB 1; Length 17;
XX Best Local Similarity 100.0%; Pred. No. 15;
XX Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
Qy 1 CTCAGCAACACTCTCTAT 17
Db 1 CTCAGCAACACTCTCTAT 17
XXXXXXXXXXXXXXXXXXXX
XXXXXXXXXXXXXXXXXXXX

RESULT 23
ADU48041
ID ADU48041 standard; DNA; 17 BP.
XX
XX AC ADU48041;
XX
XX DT 10-FEB-2005 (first entry)
XX
XX DE Human IL-1A (VNTR) polymorphic locus amplifying PCR primer #1.
XX
XX KW Restenosis; interleukin; IL; IL-1A; IL-1B; IL-1RN; vasotropic; human;
XX PCR; primer; ss.
XX
XX OS Homo sapiens.
XX
XX PN US2004229264-A1.
XX
XX PD 18-NOV-2004.
XX
XX PF 12-APR-2004; 2004US-00823197.
XX
XX PR 10-MAR-1997; 97US-00813456.
XX 26-MAY-1999; 99US-00320395.
XX 01-NOV-1999; 99US-00431352.
XX 24-MAY-2000; 2000US-00578534.
XX
XX PA (INTE-) INTERLEUKIN GENETICS INC.
XX
XX PI Crossman DC, Duff GW, Francis SE, Kornman KS, Stephenson K;
XX WPI; 2004-813241/80.
XX
XX PT New kit comprises a first primer oligonucleotide that hybridizes 5' or 3'
XX to an allele 1 of IL-1A (+4845), IL-1B (-511), IL-1B (+3954), IL-1RN
XX (VNTR), or IL-1RN (+2018), useful for diagnosing, treating, or preventing
XX restenosis.
XX

```

PS Claim 11; SEQ ID NO 7; 81bp; English.

XX The invention relates to a kit for determining the existence of or a

CC susceptibility to developing a restenosis in a subject. The kit comprises

CC of a primer that hybridizes 5' or 3' to an allele selected from allele 1

CC of the following markers, interleukin (IL)-1A (+4845), IL-1B (-511), IL-

CC 1B (+3954), IL-1RN (VNTR) or IL-1RN (+2018) or an allele in linkage

CC disequilibrium with it. The kit is useful for determining the existence

CC of or a susceptibility to developing a restenosis in a subject. It is

CC also useful for determining whether a subject has or is predisposed to

CC developing restenosis or for treating or preventing restenosis. The

CC present sequence is a PCR primer used for amplifying human IL-1A gene

CC polymorphic locus.

XX

SQ Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;

Query Match 100.0%; Score 17; DB 1; Length 17;

Best Local Similarity 100.0%; Pred. No. 15;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17

Db 1 CTCAGCAACACTCCTAT 17

RESULT 24

ADY72754

ID ADY72754 standard; DNA; 17 BP.

XX

AC ADY72754;

XX

DT 02-JUN-2005 (first entry)

DE Human IL-1RN PCR primer SEQ ID NO:5.

XX

XX interleukin-1 receptor; menopause; gynecological; gynecology; obstetrics;

KW metabolic disorder; PCR; primer; ss.

XX

OS Homo sapiens.

OS Synthetic.

XX

PN US2005064453-A1.

XX

PD 24-MAR-2005.

XX

PF 03-MAY-2004; 2004US-00838503.

XX

PR 30-JUN-1999; 99US-00345217.

PR 04-AUG-2000; 2000US-00632657.

XX

XX (DUFF/) DUFF G.

PA (KORN/) KORNMAN K.

PA (DIJK/) DIJK S V.

XX

PI Duff G, Kornman K, Dijk SV;

XX

DR WPI; 2005-241272/25.

XX

PT New kit for determining a woman's predisposition to early-onset menopause

PT (EOM) comprises primers that hybridize to a marker of an IL-1-related

PT gene associated with EOM, a DNA sampling means, a control, and a DNA

PT detection means.

XX

PS Claim 5; SEQ ID NO 5; 59pp; English.

XX

XX The invention relates to a kit for determining a woman's predisposition

CC to early-onset menopause (EOM). The kit comprises a first primer that

CC hybridizes 5' or 3' to a marker of an IL-1-related gene associated with

CC early-onset menopause. Gynecological. The kit is useful for detecting a

CC predisposition to EOM, for treating or preventing the development of EOM

CC in a woman, or for identifying agents that may treat or prevent EOM. The

CC present sequence represents a PCR primer for the human interleukin 1

CC receptor antagonist (IL-1RN) gene, which is used in an example from the

CC

CC present invention for determining an individual's allelic pattern

CC (genotyping). The human IL-1RN gene is located on chromosome 2, more

CC specifically to q13-14.1. Note: the present sequence is designated SEQ ID

CC NO:5 in the examples and claims, but corresponds with SEQ ID NO:9 in the

CC Sequence Listing.

XX

SQ Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;

Query Match 100.0%; Score 17; DB 1; Length 17;

Best Local Similarity 100.0%; Pred. No. 15;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17

Db 1 CTCAGCAACACTCCTAT 17

RESULT 25

AEE01079

ID AEE01079 standard; DNA; 17 BP.

XX

AC AEE01079;

XX

DT 26-JAN-2006 (first entry)

DE Human interleukin (IL)-1 PCR primer SEQ ID NO 22.

XX

XX periodontal disease; interleukin; IL-1; PCR; primer; ss.

XX

OS Homo sapiens.

PN WO2005108619-A2.

XX

PD 17-NOV-2005.

XX

PF 03-MAY-2005; 2005WO-US015267.

XX

PR 03-MAY-2004; 2004US-0567727P.

XX

XX (INTE-) INTERLEUKIN GENETICS INC.

XX

PI Duff GW, Kornman K, Wilkins L, Chen HM, Rogus J;

XX

DR WPI; 2005-811015/82.

XX

PT Identifying a subject having increased IL-1B transcription, for

PT identifying if the subject is predisposed to periodontal disease, by

PT providing a biological sample comprising genomic DNA, and identifying an

PT IL-1B allele in the sample.

XX

PS Example 1; SEQ ID NO 22; 126pp; English.

XX

XX The invention relates to a method of identifying a subject having

CC increased IL-1B transcription providing a biological sample comprising

CC genomic DNA from the subject, and identifying an IL-1B allele selected

CC from IL-1B (-3737) allele 1 and/or IL-1B (-1468) allele 1 in the sample,

CC where the presence of the allele indicates that the subject has increased

CC IL-1B transcription. The methods are useful for identifying if the

CC subject is predisposed to periodontal disease, and as assays for

CC identifying therapeutics for treating and/or preventing the development

CC of the disease. The present sequence represents a human interleukin (IL)-

CC 1 PCR primer.

XX

SQ Sequence 17 BP; 5 A; 7 C; 1 G; 4 T; 0 U; 0 Other;

Query Match 100.0%; Score 17; DB 1; Length 17;

Best Local Similarity 100.0%; Pred. No. 15;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17

Db 1 CTCAGCAACACTCCTAT 17

```

RESULT 26
ID ACC44794 standard; DNA; 17 BP.
XX
AC ACC44794;
XX
DT 03-JUN-2003 (first entry)
XX
DE Interleukin 1 beta PCR primer SEQ ID NO:5.
XX
KW Interleukin 1 beta; IL-1 beta; hepatitis C virus; HCV; infection;
XX liver cancer; cytostatic; PCR primer; ss.
XX
OS Homo sapiens.
XX
OS Synthetic.
XX
PN WO2003016570-A1.
XX
PD 27-FEB-2003.
XX
PF 21-AUG-2002; 2002WO-JP008415.
XX
PR 21-AUG-2001; 2001JP-00250545.
XX
PA (HUBI-) HUBIT GENOMIX INC.
XX
XX Omata M, Kato N;
XX
XX WPI; 2003-256715/25.
XX
PT Examining liver cell cancer particularly after infection with hepatitis
PT virus by using polymorphism of IL-1beta gene, also applicable in
PT screening preventives or remedies for liver cell cancer.
XX
PS Example 2; Page 33; 55pp; Japanese.
XX
CC The present invention describes a method for examining liver cell cancer
CC caused by hepatitis virus, where the method comprises the detection of a
CC polymorphism produced at the promoter region of interleukin 1 (IL-1) beta
CC gene. IL-1 beta has cytostatic activity. The method is for examining
CC liver cell cancer particularly after infection with hepatitis C virus
CC (HCV), which can also be applied in screening drugs for the prevention or
CC treatment of liver cell cancer. The present sequence represents a PCR
CC primer for IL-1 beta, which is used in an example from the present
CC invention
XX
SQ Sequence 17 BP; 4 A; 10 C; 1 G; 2 T; 0 U; 0 Other;

Query Match 82.4%; Score 14; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 42;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCC 14
Db 4 CTCAGCAACACTCC 17

RESULT 27
ADU50588
ID ADU50588 standard; DNA; 17 BP.
XX
AC ADU50588;
XX
DT 10-FEB-2005 (first entry)
XX
DE IL-1RN VNTR polymorphism PCR primer, SEQ ID 9.
XX
KW Immunosuppressive; Gene Therapy; kidney; interleukin-1; PCR; primer; ss.
XX
OS Synthetic.
XX
PN US2004229228-A1.
XX

18-NOV-2004.
24-JUL-2003; 2003US-00626830.
25-JUL-2002; 2002US-0398986P.
(NOUN ) UNIV NORTHWESTERN.
PA (AMGE-) AMGEN INC.
XX
PI Sims JE, Kaufman DB;
XX
DR WPI; 2005-010687/01.
XX
PT Diagnosing early kidney allograft rejection comprises determining a
PT genotype of an IL-1 family member gene.
XX
PS Example 1; SEQ ID NO 9; 12pp; English.
XX
CC The present invention relates to a method for identifying an individual
CC predisposed to early rejection of a kidney allograft. The method
CC comprises determining a genotype of an interleukin-1 (IL-1) family member
CC gene, where the presence of one or more alleles, e.g. an A1,A2 allele
CC combination at an IL-1A VNTR intron 6 locus, an A2 allele at an IL-
CC 1A-4845 locus, or an E2 allele at an IL-1B-3953 locus indicates that the
CC individual is at increased risk for early rejection of the kidney
CC allograft. Also claimed is a primer set for determining a genotype in an
CC IL-1 family member gene. The present sequence is one such primer. The
CC methods and primers are useful for identifying an individual predisposed
CC to and treating a patient predisposed to early rejection of a kidney
CC allograft.
XX
SQ Sequence 17 BP; 4 A; 10 C; 1 G; 2 T; 0 U; 0 Other;

Query Match 82.4%; Score 14; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 42;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCC 14
Db 4 CTCAGCAACACTCC 17

RESULT 28
AAD48159/c
ID AAD48159 standard; DNA; 16 BP.
XX
AC AAD48159;
XX
DT 24-FEB-2003 (first entry)
XX
DE PCR primer #7 used for single nucleotide polymorphism (SNP) analysis.
XX
KW Peptide nucleic acid; PNA; nucleic acid zygosity; genetic analysis;
XX scientific investigation; pharmacogenomic; pharmacogenetic; epigenomic;
XX PCR; primer; ss.
XX
OS Unidentified.
XX
PN WO200272865-A2.
XX
PD 19-SEP-2002.
XX
PF 09-MAR-2002; 2002WO-US007050.
XX
PR 09-MAR-2001; 2001US-0274547P.
XX
PA (BOST-) BOSTON PROBES INC.
XX
PI Coull JM, Flandaca MJ, Kristjanson MD, Hyldig-Nielsen JJ;
XX Creasey TM;
XX
XX WPI; 2003-018741/01.
XX

```

XX Composition for determining target sequence of contiguous nucleobases.
 PT comprises polynucleobase strand and combination oligomer comprising first
 PT and second oligomer blocks that are covalently linked to each other.

XX Example 5; Page 70; 149pp; English.

XX The present invention relates to combination oligomers, including block
 CC synthesis of combination of oligomers in the absence of a template. The
 CC invention relates to a composition comprising a polynucleobase strand and
 CC a combination oligomer comprising first and second oligomer blocks that
 CC are each independently a peptide nucleic acid (PNA) covalently linked to
 CC each other by a linker of at least three atoms in length, where the
 CC oligomer blocks are sequences specifically hybridised to a target
 CC sequence of contiguous nucleobases in the polynucleobase strand. To form
 CC a double stranded target sequence-oligomer complex. The composition is
 CC used for determining a target sequence of contiguous nucleobases and for
 CC determining the zygosity of a nucleic acid for a single nucleotide
 CC polymorphism (SNP). The methods are useful in scientific investigation,
 CC e.g., for detection, identification and/or enumeration of bacteria,
 CC viruses and pathogens in food, beverages, water, pharmaceutical products,
 CC personal care products, dairy products, in clinical samples or in samples
 CC of plant, animal, human or environmental origin. They are also useful for
 CC the analysis of raw materials, equipment, products or processes used to
 CC manufacture or store food, beverages, water, pharmaceutical products,
 CC personal care products dairy products or environmental samples. The
 CC methods and materials are useful in areas such as expression analysis,
 CC SNP analysis, genetic analysis of humans, animals, fungi, yeast viruses
 CC and plants, therapy monitoring, pharmacogenomics, pharmacogenetics,
 CC epigenomics and high throughput screening operations. The present
 CC sequence is a PCR primer used for single nucleotide polymorphism (SNP)
 CC analysis

XX SQ Sequence 16 BP; 3 A; 1 C; 8 G; 4 T; 0 U; 0 Other;

Query Match 72.9%; Score 12.4; DB 1; Length 16;
 Best Local Similarity 92.9%; Pred. No. 68;
 Matches 13; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 TCAGCAACACTCTCT 15

DB 15 TCAGCAACACTCTCT 2

RESULT 29

ID ACN01682
 ACN01682 standard; RNA; 17 BP.

XX ACN01682;

XX 22-APR-2004 (first entry)

XX WNV Inozyme substrate SEQ ID NO 1672.

XX WNV; West Nile Virus; antiinflammatory; cytostatic; hepatotropic;
 KW virucide; neuroprotective; antibacterial; replication; pancreatitis;
 KW encephalitis; myocarditis; meningitis; infection; hepatitis;
 KW liver failure; cancer; cirrhosis; Hammerhead; Inozyme; DNAzyme;
 KW Amberzyme; Zinzyne; ss.

XX West Nile Virus.

XX WO200268637-A2.

XX 06-SEP-2002.

XX 19-OCT-2001; 2001WO-US048350.

XX 20-OCT-2000; 2000US-0242411P.

XX (RIBO-) RIBOZYME PHARM INC.

XX (BLAT/) BLATT L.

XX (MCSW/) MCSWIGGEN J A.

XX Blatt L, Mcswiggen JA;

XX WPI; 2002-706994/76.

XX New nucleic acid molecule that modulates replication of West Nile Virus
 PT (WNV), useful for treating a condition related to WNV infection e.g.
 PT pancreatitis, meningitis, hepatocellular carcinoma or cirrhosis.

XX Claim 23; SEQ ID NO 1672; 495pp; English.

XX The invention relates to nucleic acid molecules that modulate replication
 CC of the West Nile Virus (WNV). The nucleic acid molecules are useful for
 CC treating a condition related to WNV infection e.g. pancreatitis,
 CC encephalitis, myocarditis, meningitis, neurologic infection, hepatitis,
 CC liver failure, hepatocellular carcinoma or cirrhosis. The nucleic acid
 CC molecule is selected from the group of ribozymes consisting of
 CC Hammerhead, inozyme, G-cleaver, DNAzyme, Amberzyme and zinzyne. The
 CC nucleic acid molecules further comprise at least five ribose residues, at
 CC least ten 2'-O-methyl modifications, phosphorothioate linkages on at
 CC least three of the 5' terminal nucleotides and a 3' end modification of a
 CC 3'-3' inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080
 CC are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given
 CC in the specification. The present sequence is that of a nucleic acid
 CC molecule of the invention

XX SQ Sequence 17 BP; 4 A; 6 C; 2 G; 0 T; 5 U; 0 Other;

Query Match 72.9%; Score 12.4; DB 1; Length 17;

Best Local Similarity 71.4%; Pred. No. 74;

Matches 10; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 2 TCAGCAACACTCTCT 15

DB 4 UCAGCAUCACUCCU 17

RESULT 30

ID ACN01683
 ACN01683 standard; RNA; 17 BP.

XX ACN01683;

XX 22-APR-2004 (first entry)

XX WNV Inozyme substrate SEQ ID NO 1673.

XX WNV; West Nile Virus; antiinflammatory; cytostatic; hepatotropic;
 KW virucide; neuroprotective; antibacterial; replication; pancreatitis;
 KW encephalitis; myocarditis; meningitis; infection; hepatitis;
 KW liver failure; cancer; cirrhosis; Hammerhead; Inozyme; DNAzyme;
 KW Amberzyme; Zinzyne; ss.

XX West Nile Virus.

XX WO200268637-A2.

XX 06-SEP-2002.

XX 19-OCT-2001; 2001WO-US048350.

XX 20-OCT-2000; 2000US-0242411P.

XX (RIBO-) RIBOZYME PHARM INC.

XX (BLAT/) BLATT L.

XX (MCSW/) MCSWIGGEN J A.

XX Blatt L, Mcswiggen JA;

XX WPI; 2002-706994/76.

XX New nucleic acid molecule that modulates replication of West Nile Virus
 PT (WNV), useful for treating a condition related to WNV infection e.g.

```

PT pancreatitis, meningitis, hepatocellular carcinoma or cirrhosis.
XX Claim 23; SEQ ID NO 1673; 495pp; English.
PS
XX
CC The invention relates to nucleic acid molecules that modulate replication
CC of the West Nile Virus (WNV). The nucleic acid molecules are useful for
CC treating a condition related to WNV infection e.g. pancreatitis,
CC encephalitis, myocarditis, meningitis, neurologic infection, hepatitis,
CC liver failure, hepatocellular carcinoma or cirrhosis. The nucleic acid
CC molecule is selected from the group of ribozymes consisting of
CC Hammerhead, Inozyme, G-cleaver, DNazyme, Amberzyme and Zinzyme. The
CC nucleic acid molecules further comprise at least five ribose residues, at
CC least ten 2'-O-methyl modifications, phosphorothioate linkages on at
CC least three of the 5' terminal nucleotides and a 3' end modification of a
CC 3'-3' inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080
CC are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given
CC in the specification. The present sequence is that of a nucleic acid
CC molecule of the invention
XX
SQ Sequence 17 BP; 3 A; 7 C; 3 G; 0 T; 4 U; 0 Other;

Query Match 72.9%; Score 12.4; DB 1; Length 17;
Best Local Similarity 71.4%; Pred. NO. 74;
Matches 10; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Qy 2 TCAGCAACACTCTCT 15
Db 1 UCAGCAUCACUCCU 14

RESULT 31
ACN00198
ID ACN00198 standard; RNA; 17 BP.
XX
AC ACN00198;
XX
XX 22-APR-2004 (first entry)
XX
DE WNV Hammerhead Ribozyme substrate SEQ ID NO 188.
XX
XX WNV, West Nile Virus; antiinflammatory; cytosolic; hepatotropic;
XX virucide; neuroprotective; antibacterial; replication; pancreatitis;
XX encephalitis; myocarditis; meningitis; infection; hepatitis;
XX liver failure; cancer; cirrhosis; Hammerhead; Inozyme; DNazyme;
XX Amberzyme; Zinzyme; ss.
XX
OS West Nile Virus.
XX
XX WO200268637-A2.
XX
XX 06-SEP-2002.
XX
XX 19-OCT-2001; 2001WO-US048350.
XX
XX 20-OCT-2000; 2000US-024241P.
XX
XX (RIBO-) RIBOZYME PHARM INC.
XX (BLAT/) BLATT L.
XX (MCSW/) MCSWIGGEN J A.
XX
XX Blatt L, Mcswiggen JA;
XX
XX WPI; 2002-706994/76.
XX
XX New nucleic acid molecule that modulates replication of West Nile Virus
XX (WNV), useful for treating a condition related to WNV infection e.g.
XX pancreatitis, meningitis, hepatocellular carcinoma or cirrhosis.
XX
XX Claim 23; SEQ ID NO 188; 495pp; English.
XX
XX The invention relates to nucleic acid molecules that modulate replication
XX of the West Nile Virus (WNV). The nucleic acid molecules are useful for
XX treating a condition related to WNV infection e.g. pancreatitis,
XX encephalitis, myocarditis, meningitis, neurologic infection, hepatitis,
XX liver failure, hepatocellular carcinoma or cirrhosis. The nucleic acid
XX molecule is selected from the group of ribozymes consisting of
XX Hammerhead, Inozyme, G-cleaver, DNazyme, Amberzyme and Zinzyme. The
XX nucleic acid molecules further comprise at least five ribose residues, at
XX least ten 2'-O-methyl modifications, phosphorothioate linkages on at
XX least three of the 5' terminal nucleotides and a 3' end modification of a
XX 3'-3' inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080
XX are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given
XX in the specification. The present sequence is that of a nucleic acid
XX molecule of the invention
XX
SQ Sequence 17 BP; 3 A; 7 C; 3 G; 0 T; 4 U; 0 Other;

Query Match 72.9%; Score 12.4; DB 1; Length 17;
Best Local Similarity 71.4%; Pred. NO. 74;
Matches 10; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Qy 2 TCAGCAACACTCTCT 15
Db 1 UCAGCAUCACUCCU 14

RESULT 32
ACN14095/C
ID ACN14095 standard; RNA; 17 BP.
XX
AC ACN14095;
XX
XX 22-APR-2004 (first entry)
XX
DE WNV minus strand DNazyme substrate SEQ ID NO 14098.
XX
XX WNV, West Nile Virus; antiinflammatory; cytosolic; hepatotropic;
XX virucide; neuroprotective; antibacterial; replication; pancreatitis;
XX encephalitis; myocarditis; meningitis; infection; hepatitis;
XX liver failure; cancer; cirrhosis; Hammerhead; Inozyme; DNazyme;
XX Amberzyme; Zinzyme; ss.
XX
OS West Nile Virus.
XX
XX WO200268637-A2.
XX
XX 06-SEP-2002.
XX
XX 19-OCT-2001; 2001WO-US048350.
XX
XX 20-OCT-2000; 2000US-024241P.
XX
XX (RIBO-) RIBOZYME PHARM INC.
XX (BLAT/) BLATT L.
XX (MCSW/) MCSWIGGEN J A.
XX
XX Blatt L, Mcswiggen JA;
XX
XX WPI; 2002-706994/76.
XX
XX New nucleic acid molecule that modulates replication of West Nile Virus
XX (WNV), useful for treating a condition related to WNV infection e.g.
XX pancreatitis, meningitis, hepatocellular carcinoma or cirrhosis.
XX
XX Claim 23; SEQ ID NO 14098; 495pp; English.
XX
XX The invention relates to nucleic acid molecules that modulate replication
XX of the West Nile Virus (WNV). The nucleic acid molecules are useful for
XX treating a condition related to WNV infection e.g. pancreatitis,
XX encephalitis, myocarditis, meningitis, neurologic infection, hepatitis,
XX liver failure, hepatocellular carcinoma or cirrhosis. The nucleic acid
XX molecule is selected from the group of ribozymes consisting of
XX Hammerhead, Inozyme, G-cleaver, DNazyme, Amberzyme and Zinzyme. The
XX nucleic acid molecules further comprise at least five ribose residues, at
XX least ten 2'-O-methyl modifications, phosphorothioate linkages on at
XX least three of the 5' terminal nucleotides and a 3' end modification of a
XX 3'-3' inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080
XX are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given
XX in the specification. The present sequence is that of a nucleic acid
XX molecule of the invention
XX
SQ Sequence 17 BP; 3 A; 7 C; 2 G; 0 T; 5 U; 0 Other;

Query Match 72.9%; Score 12.4; DB 1; Length 17;
Best Local Similarity 71.4%; Pred. NO. 74;
Matches 10; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Qy 2 TCAGCAACACTCTCT 15
Db 2 UCAGCAUCACUCCU 15

RESULT 33
ACN14095/C
ID ACN14095 standard; RNA; 17 BP.
XX
AC ACN14095;
XX
XX 22-APR-2004 (first entry)
XX
DE WNV minus strand DNazyme substrate SEQ ID NO 14098.
XX
XX WNV, West Nile Virus; antiinflammatory; cytosolic; hepatotropic;
XX virucide; neuroprotective; antibacterial; replication; pancreatitis;
XX encephalitis; myocarditis; meningitis; infection; hepatitis;
XX liver failure; cancer; cirrhosis; Hammerhead; Inozyme; DNazyme;
XX Amberzyme; Zinzyme; ss.
XX
OS West Nile Virus.
XX
XX WO200268637-A2.
XX
XX 06-SEP-2002.
XX
XX 19-OCT-2001; 2001WO-US048350.
XX
XX 20-OCT-2000; 2000US-024241P.
XX
XX (RIBO-) RIBOZYME PHARM INC.
XX (BLAT/) BLATT L.
XX (MCSW/) MCSWIGGEN J A.
XX
XX Blatt L, Mcswiggen JA;
XX
XX WPI; 2002-706994/76.
XX
XX New nucleic acid molecule that modulates replication of West Nile Virus
XX (WNV), useful for treating a condition related to WNV infection e.g.
XX pancreatitis, meningitis, hepatocellular carcinoma or cirrhosis.
XX
XX Claim 23; SEQ ID NO 14098; 495pp; English.
XX
XX The invention relates to nucleic acid molecules that modulate replication
XX of the West Nile Virus (WNV). The nucleic acid molecules are useful for
XX treating a condition related to WNV infection e.g. pancreatitis,
XX encephalitis, myocarditis, meningitis, neurologic infection, hepatitis,
XX liver failure, hepatocellular carcinoma or cirrhosis. The nucleic acid
XX molecule is selected from the group of ribozymes consisting of
XX Hammerhead, Inozyme, G-cleaver, DNazyme, Amberzyme and Zinzyme. The
XX nucleic acid molecules further comprise at least five ribose residues, at
XX least ten 2'-O-methyl modifications, phosphorothioate linkages on at
XX least three of the 5' terminal nucleotides and a 3' end modification of a
XX 3'-3' inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080
XX are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given
XX in the specification. The present sequence is that of a nucleic acid
XX molecule of the invention
XX
SQ Sequence 17 BP; 3 A; 7 C; 2 G; 0 T; 5 U; 0 Other;

Query Match 72.9%; Score 12.4; DB 1; Length 17;
Best Local Similarity 71.4%; Pred. NO. 74;
Matches 10; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Qy 2 TCAGCAACACTCTCT 15
Db 2 UCAGCAUCACUCCU 15

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CC 3'-3' inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080
 CC are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given
 CC in the specification. The present sequence is that of a nucleic acid
 CC molecule of the invention

XX Sequence 17 BP; 5 A; 2 C; 6 G; 0 T; 4 U; 0 Other;

Query Match 72.9%; Score 12.4; DB 1; Length 17;
 Best Local Similarity 92.9%; Pred. No. 74;
 Matches 13; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 TCAGCAACACTCT 15
 Db 15 TCAGCACTACTCT 2

RESULT 33
 ACN15151/c
 ID ACN15151 standard, RNA; 17 BP.

XX ACN15151;

AC ACN15151;

XX 22-APR-2004 (first entry)

DE WNV minus strand Amberzyme substrate SEQ ID NO 15154.

XX WNV; West Nile Virus; antiinflammatory; cytostatic; hepatotropic;
 KW viricide; neuroprotective; antibacterial; replication; pancreatitis;
 KW encephalitis; myocarditis; meningitis; infection; hepatitis;
 KW liver failure; cancer; cirrhosis; Hammerhead; Inozyme; DNAzyme;
 KW Amberzyme; Zinzyme; ss.

XX West Nile Virus.

XX WO200268637-A2.

XX 06-SEP-2002.

XX 19-OCT-2001; 2001WO-US048350.

XX 20-OCT-2000; 2000US-0242411P.

XX (RIBO-) RIBOZYME PHARM INC.

XX (BLAT/) BLATT L.

XX (MCSW/) MCSWIGGEN J A.

XX Blatt L, Mcswiggen JA;

XX WPI; 2002-706994/76.

XX New nucleic acid molecule that modulates replication of West Nile Virus

PT (WNV), useful for treating a condition related to WNV infection e.g.

PT pancreatitis, meningitis, hepatocellular carcinoma or cirrhosis.

XX Claim 23; SEQ ID NO 15154; 495pp; English.

XX The invention relates to nucleic acid molecules that modulate replication
 CC of the West Nile Virus (WNV). The nucleic acid molecules are useful for
 CC treating a condition related to WNV infection e.g. pancreatitis,
 CC encephalitis, myocarditis, meningitis, neurologic infection, hepatitis,
 CC liver failure, hepatocellular carcinoma or cirrhosis. The nucleic acid
 CC molecule is selected from the group of ribozymes consisting of
 CC Hammerhead, Inozyme, G-cleaver, DNAzyme, Amberzyme and Zinzyme. The
 CC nucleic acid molecules further comprise at least five ribose residues, at
 CC least ten 2'-O-methyl modifications, phosphorothioate linkages on at
 CC least three of the 5' terminal nucleotides and a 3' end modification of a
 CC 3'-3' inverted abasic moiety. Nucleic acid molecules SEQ ID NO 1 to 37080
 CC are claimed; however, SEQ ID NO 2194-2206 and 17502-17514 are not given
 CC in the specification. The present sequence is that of a nucleic acid
 CC molecule of the invention

XX Sequence 17 BP; 5 A; 2 C; 7 G; 0 T; 3 U; 0 Other;

Query Match 72.9%; Score 12.4; DB 1; Length 17;
 Best Local Similarity 92.9%; Pred. No. 74;
 Matches 13; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 TCAGCAACACTCT 15
 Db 16 TCAGCACTACTCT 3

RESULT 34
 ABI02395/c
 ID ABI02395 standard; DNA; 12 BP.

XX ABI02395;

XX 22-FEB-2002 (first entry)

XX Oligonucleotide primer SEQ ID NO 302368 for detecting SNP TSC0019966.

XX SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
 KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
 KW central nervous system; gastrointestinal; respiratory; immune; metabolic.

XX Homo sapiens.

XX WO200177384-A2.

XX 18-OCT-2001.

XX 06-APR-2001; 2001WO-IB000713.

XX 07-APR-2000; 2000DE-01019173.

XX (EPIG-) EPIGENOMICS AG.

XX Olek A, Piepenbrock C, Berlin K;

XX WPI; 2001-657177/75.

XX Set of oligonucleotides, useful for diagnosis and cell typing, is
 PT designed to detect single-nucleotide polymorphisms and cytosine
 PT methylation status.

XX Claim 1; SEQ ID NO 302368; 29pp + Sequence Listing; German.

XX This invention describes novel oligonucleotide primers or peptide nucleic
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
 CC and cytosine methylation status in chemically pretreated genomic DNA. The
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a
 CC range of diseases including immune system, gastrointestinal, respiratory,
 CC central nervous system, cardiovascular and metabolic disorders. The
 CC oligomers are also used for detecting cell type differentiation. ABC00010
 CC -ABC9989, ABF0010-ABF9989, ABH0010-ABH9989 and ABI0010-ABI82073
 CC represent the oligomers described in the invention. NOTE: The sequence
 CC data for this patent did not form part of the printed specification, but
 CC was obtained in electronic format from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences

XX Sequence 12 BP; 3 A; 0 C; 5 G; 4 T; 0 U; 0 Other;

Query Match 64.7%; Score 11; DB 1; Length 12;
 Best Local Similarity 100.0%; Pred. No. 76;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 6 CAACACTCTCTA 16
 Db 11 CAACACTCTCTA 1

RESULT 35
 ABK72568
 ID ABK72568 standard; DNA; 12 BP.

XX

```

AC ABK72568;
XX
DT 13-AUG-2002 (first entry)
XX
DE Human OPAL gene, exon/intron junction #35.
XX
KW Human; ophthalmological; OPAL; autosomal dominant optic atrophy; ADOA;
KW gene; ds.
XX
OS Homo sapiens.
XX
XX WO200227022-A2.
XX
PD 04-APR-2002.
XX
PF 26-SEP-2001; 2001WO-GB004284.
XX
PR 26-SEP-2000; 2000GB-00023555.
XX
XX (UNLO ) UNIV COLLEGE LONDON.
PA (UYEY-) UNIV EYE HOSPITAL.
XX
PI Bhattacharya S, Wisinger B, Alexander C, Votruba M;
XX
DR WPI; 2002-416484/44.
XX
XX Novel human normal or mutant OPAL (the predominant locus for autosomal
PT dominant optic atrophy (ADOA)) polypeptides and the OPAL gene, useful in
PT the diagnosis and treatment of autosomal dominant optic atrophy ADOA.
XX
XX Disclosure; Fig 12; 75pp; English.
XX
XX The invention relates to an isolated human normal or mutant OPAL (the
CC predominant locus for autosomal dominant optic atrophy (ADOA))
CC polypeptide (I), characterised by a molecular weight of about 112 kDa,
CC and substantially free of other human proteins. Also described is the DNA
CC (ii) encoding (i). (i) and (ii) are useful as a medicament, for the
CC treatment of a medical condition resulting from a defect in the OPAL
CC gene, which results in autosomal dominant optic atrophy. The nucleic acid
CC and antibodies to (i) are useful in a variety of hybridisation and
CC immunological assays to screen for, and to detect the presence of, either
CC a normal or a defective OPAL gene or gene product. ABK72533-ABK72593
CC represent the human OPAL gene and intron/exon splice junctions
XX
SQ Sequence 12 BP; 4 A; 6 C; 1 G; 1 T; 0 U; 0 Other;

Query Match 64.7%; Score 11; DB 1; Length 12;
Best Local Similarity 100.0%; Pred. No. 76;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACAC 11
Db 2 CTCAGCAACAC 12
|||||
|||||

RESULT 36
ABF36021
ID ABF36021 standard; DNA; 13 BP.
XX
AC ABF36021;
XX
DT 21-FEB-2002 (first entry)
XX
DE Oligonucleotide SEQ ID NO 136018 for detecting SNP TSC0033969.
XX
XX SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
XX
OS Homo sapiens.
XX
XX WO200177384-A2.
XX
PN WO200177384-A2.
XX

Query Match 64.7%; Score 11; DB 1; Length 13;
Best Local Similarity 100.0%; Pred. No. 84;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 7 AACACTCCTAT 17
Db 2 AACACTCCTAT 12
|||||
|||||

RESULT 37
ABC71077
ID ABC71077 standard; DNA; 13 BP.
XX
AC ABC71077;
XX
DT 21-FEB-2002 (first entry)
XX
DE Oligonucleotide SEQ ID NO 71094 for detecting SNP TSC0018437.
XX
XX SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
XX
OS Homo sapiens.
XX
XX WO200177384-A2.
XX
PN WO200177384-A2.
XX
PD 18-OCT-2001.
XX
PF 06-APR-2001; 2001WO-IB000713.
XX
PR 07-APR-2000; 2000DE-01019173.
XX
XX (EPIG-) EPIGENOMICS AG.
PA
PI Olek A, Piepenbrock C, Berlin K;
XX
DR WPI; 2001-657177/75.
XX
XX Set of oligonucleotides, useful for diagnosis and cell typing, is
PT designed to detect single-nucleotide polymorphisms and cytosine
PT methylation status.
XX
PS Claim 1; SEQ ID NO 136018; 29pp + Sequence Listing; German.
XX
XX This invention describes novel oligonucleotide primers or peptide nucleic
CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
CC and cytosine methylation status in chemically pretreated genomic DNA. The
CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a
CC range of diseases including immune system, gastrointestinal, respiratory,
CC central nervous system, cardiovascular and metabolic disorders. The
CC oligomers are also used for detecting cell type differentiation. ABC00010
CC -ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and ABI00010-ABI82073
CC represent the oligomers described in the invention. NOTE: The sequence
CC data for this patent did not form part of the printed specification, but
CC was obtained in electronic format from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences
XX
SQ Sequence 13 BP; 6 A; 4 C; 0 G; 3 T; 0 U; 0 Other;

Query Match 64.7%; Score 11; DB 1; Length 13;
Best Local Similarity 100.0%; Pred. No. 84;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 7 AACACTCCTAT 17
Db 2 AACACTCCTAT 12
|||||
|||||

RESULT 37
ABC71077
ID ABC71077 standard; DNA; 13 BP.
XX
AC ABC71077;
XX
DT 21-FEB-2002 (first entry)
XX
DE Oligonucleotide SEQ ID NO 71094 for detecting SNP TSC0018437.
XX
XX SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
XX
OS Homo sapiens.
XX
XX WO200177384-A2.
XX
PN WO200177384-A2.
XX
PD 18-OCT-2001.
XX
PF 06-APR-2001; 2001WO-IB000713.
XX
PR 07-APR-2000; 2000DE-01019173.
XX
XX (EPIG-) EPIGENOMICS AG.
PA
PI Olek A, Piepenbrock C, Berlin K;
XX
DR WPI; 2001-657177/75.
XX
XX Set of oligonucleotides, useful for diagnosis and cell typing, is
PT designed to detect single-nucleotide polymorphisms and cytosine
PT methylation status.

```


PT designed to detect single-nucleotide polymorphisms and cytosine
 XX methylation status.
 XX
 PS Claim 1; SEQ ID NO 71094; 29pp + Sequence Listing; German.
 XX
 CC This invention describes novel oligonucleotide primers or peptide nucleic
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
 CC and cytosine methylation status in chemically pretreated genomic DNA. The
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a
 CC range of diseases including immune system, gastrointestinal, respiratory,
 CC central nervous system, cardiovascular and metabolic disorders. The
 CC oligomers are also used for detecting cell type differentiation. ABC00010
 CC -ABCF99989, ABF00010-ABF99989, ABH00010-ABH99989 and ABI00010-ABI82073
 CC represent the oligomers described in the invention. NOTE: The sequence
 CC data for this patent did not form part of the printed specification, but
 CC was obtained in electronic format from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX
 XX Sequence 13 BP; 4 A; 6 C; 0 G; 3 T; 0 U; 0 Other;
 SQ
 Query Match 64.7%; Score 11; DB 1; Length 13;
 Best Local Similarity 100.0%; Pred. No. 84;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 6 CAACACTCCTTA 16
 Db 1 CAACACTCCTTA 11
 |||||
 |||||
 RESULT 38
 ABF36020/c
 ID ABF36020 standard; DNA; 13 BP.
 XX
 AC ABF36020;
 XX
 DT 21-FEB-2002 (first entry)
 XX
 DE Oligonucleotide SEQ ID NO 136017 for detecting SNP TSC0033969.
 XX
 KW SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
 KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
 KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
 XX
 OS Homo sapiens.
 XX
 PN WO200177384-A2.
 XX
 PD 18-OCT-2001.
 XX
 PF 06-APR-2001; 2001WO-IB000713.
 XX
 PR 07-APR-2000; 2000DE-01019173.
 XX
 PA (EPIG-) EPIGENOMICS AG.
 XX
 PI Olek A, Piepenbrock C, Berlin K;
 XX
 DR WPI; 2001-657177/75.
 XX
 XX Set of oligonucleotides, useful for diagnosis and cell typing, is
 PT designed to detect single-nucleotide polymorphisms and cytosine
 PT methylation status.
 XX
 PS Claim 1; SEQ ID NO 136017; 29pp + Sequence Listing; German.
 XX
 CC This invention describes novel oligonucleotide primers or peptide nucleic
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
 CC and cytosine methylation status in chemically pretreated genomic DNA. The
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a
 CC range of diseases including immune system, gastrointestinal, respiratory,
 CC central nervous system, cardiovascular and metabolic disorders. The
 CC oligomers are also used for detecting cell type differentiation. ABC00010
 CC -ABCF99989, ABF00010-ABF99989, ABH00010-ABH99989 and ABI00010-ABI82073

CC represent the oligomers described in the invention. NOTE: The sequence
 CC data for this patent did not form part of the printed specification, but
 CC was obtained in electronic format from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX
 XX Sequence 13 BP; 3 A; 0 C; 4 G; 6 T; 0 U; 0 Other;
 SQ
 Query Match 64.7%; Score 11; DB 1; Length 13;
 Best Local Similarity 100.0%; Pred. No. 84;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 7 AACACTCCTAT 17
 Db 12 AACACTCCTAT 2
 |||||
 |||||
 RESULT 39
 ABC71076/c
 ID ABC71076 standard; DNA; 13 BP.
 XX
 AC ABC71076;
 XX
 DT 21-FEB-2002 (first entry)
 XX
 DE Oligonucleotide SEQ ID NO 71093 for detecting SNP TSC0018437.
 XX
 KW SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
 KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
 KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
 XX
 OS Homo sapiens.
 XX
 PN WO200177384-A2.
 XX
 PD 18-OCT-2001.
 XX
 PF 06-APR-2001; 2001WO-IB000713.
 XX
 PR 07-APR-2000; 2000DE-01019173.
 XX
 PA (EPIG-) EPIGENOMICS AG.
 XX
 PI Olek A, Piepenbrock C, Berlin K;
 XX
 DR WPI; 2001-657177/75.
 XX
 XX Set of oligonucleotides, useful for diagnosis and cell typing, is
 PT designed to detect single-nucleotide polymorphisms and cytosine
 PT methylation status.
 XX
 PS Claim 1; SEQ ID NO 71093; 29pp + Sequence Listing; German.
 XX
 CC This invention describes novel oligonucleotide primers or peptide nucleic
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
 CC and cytosine methylation status in chemically pretreated genomic DNA. The
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a
 CC range of diseases including immune system, gastrointestinal, respiratory,
 CC central nervous system, cardiovascular and metabolic disorders. The
 CC oligomers are also used for detecting cell type differentiation. ABC00010
 CC -ABCF99989, ABF00010-ABF99989, ABH00010-ABH99989 and ABI00010-ABI82073
 CC represent the oligomers described in the invention. NOTE: The sequence
 CC data for this patent did not form part of the printed specification, but
 CC was obtained in electronic format from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX
 XX Sequence 13 BP; 3 A; 0 C; 6 G; 4 T; 0 U; 0 Other;
 SQ
 Query Match 64.7%; Score 11; DB 1; Length 13;
 Best Local Similarity 100.0%; Pred. No. 84;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 6 CAACACTCCTTA 16
 |||||
 |||||

Db 13 CAACACTCCTTA 3

RESULT 40

ABX03942/c

ID ABX03942 standard; DNA; 15 BP.

XX AC ABX03942;

XX DT 09-JAN-2003 (first entry)

XX DE S. cerevisiae 28S rRNA fragment.

XX KW Detection; probe; diagnosis; oral disease; parodontitis; caries; therapy;

XX KW polymorphism; virulence factor; antibiotic resistance gene; prognosis;

XX KW oral infection; detection; pathogen; coronary heart disease;

XX KW diabetic symptom; ss.

XX OS Saccharomyces cerevisiae.

XX PN DE20110013-UI.

XX PD 18-OCT-2001.

XX PF 13-MAR-2001; 2001DE-02010013.

XX PR 13-MAR-2001; 2001DE-01012348.

XX PR 13-MAR-2001; 2001DE-02010013.

XX PA (ROET/) ROETGER A.

XX PA WPI; 2001-657777/76.

XX PT Oligonucleotide array, useful for diagnosing oral diseases, particularly

XX PT parodontitis, carries human or microbial reference sequences.

XX PS Claim 8; Page 22; 58pp; German.

XX SQ This invention describes a novel nucleotide carrier with probes used for

CC diagnosis of oral diseases, particularly parodontitis, but also caries,

CC especially to identify genetic predisposition (as indicated by

CC polymorphisms) to disease and to identify causative microorganisms or

CC their associated virulence factors and antibiotic resistance genes, e.g.

CC for selection of therapy and for prognosis. They are also useful for

CC research into oral infections. The carriers allow simultaneous detection

CC of both host and pathogen parameters, providing quickly and simply an

CC individual's parodontitis profile, including detection of pathogens that

CC are associated with increased risk of coronary heart diseases and/or

CC aggravation of diabetic symptoms, and of opportunistic pathogens.

CC ABX03870-ABX04044 represent DNA fragments used to illustrate the method

CC of the invention

XX SQ Sequence 15 BP; 5 A; 2 C; 4 G; 4 T; 0 U; 0 Other;

Query Match 63.5%; Score 10.8; DB 1; Length 15;

Best Local Similarity 85.7%; Pred. No. 1.1e+02;

Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 4 AGCACTCCTAT 17

DB 15 AGCTACATTCCTAT 2

RESULT 41

ABH90683/c

ID ABH90683 standard; DNA; 12 BP.

XX AC ABH90683;

XX DT 22-FEB-2002 (first entry)

XX DE Oligonucleotide primer SEQ ID NO 290676 for detecting SNP TSC0014465.

XX

KW SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;

KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;

KW central nervous system; gastrointestinal; respiratory; immune; metabolic.

OS Homo sapiens.

XX WO200177384-A2.

XX PD 18-OCT-2001.

XX PF 06-APR-2001; 2001WO-IB000713.

XX PR 07-APR-2000; 2000DE-01019173.

XX PA (SPIG-) EPIGENOMICS AG.

XX PI Olek A, Piepenbrock C, Berlin K;

XX DR WPI; 2001-657177/75.

XX PT Set of oligonucleotides, useful for diagnosis and cell typing, is

PT designed to detect single-nucleotide polymorphisms and cytosine

PT methylation status.

XX Claim 1; SEQ ID NO 290676; 29pp + Sequence Listing; German.

XX This invention describes novel oligonucleotide primers or peptide nucleic

CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)

CC and cytosine methylation status in chemically pretreated genomic DNA. The

CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a

CC range of diseases including immune system, gastrointestinal, respiratory,

CC central nervous system, cardiovascular and metabolic disorders. The

CC oligomers are also used for detecting cell type differentiation. ABC00010

CC -ABC9989, ABF00010-ABF9989, ABH00010-ABH9989 and ABT00010-ABT82073

CC represent the oligomers described in the invention. NOTE: The sequence

CC data for this patent did not form part of the printed specification, but

CC was obtained in electronic format from WIPO at

CC ftp.wipo.int/pub/published_pct_sequences

XX SQ Sequence 12 BP; 3 A; 0 C; 4 G; 5 T; 0 U; 0 Other;

Query Match 61.2%; Score 10.4; DB 1; Length 12;

Best Local Similarity 91.7%; Pred. No. 93;

Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 6 CAACACTCCTAT 17

DB 12 CAACACTCCTAT 1

RESULT 42

ABI59938/c

ID ABI59938 standard; DNA; 12 BP.

XX AC ABI59938;

XX DT 22-FEB-2002 (first entry)

XX DE Oligonucleotide primer SEQ ID NO 359911 for detecting SNP TSC0051838.

XX KW SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;

XX KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;

XX KW central nervous system; gastrointestinal; respiratory; immune; metabolic.

XX OS Homo sapiens.

XX WO200177384-A2.

XX PD 18-OCT-2001.

XX PF 06-APR-2001; 2001WO-IB000713.

XX PR 07-APR-2000; 2000DE-01019173.

XX (EPIG-) EPIGENOMICS AG.
 PA Olek A, Piepenbrock C, Berlin K;
 PI WPI; 2001-657177/75.
 DR
 XX
 XX Set of oligonucleotides, useful for diagnosis and cell typing, is
 PT designed to detect single-nucleotide polymorphisms and cytosine
 PT methylation status.
 XX
 XX Claim 1; SEQ ID NO 359911; 29pp + Sequence Listing; German.
 XX
 XX This invention describes novel oligonucleotide primers or peptide nucleic
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
 CC and cytosine methylation status in chemically pretreated genomic DNA. The
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a
 CC range of diseases including immune system, gastrointestinal, respiratory,
 CC central nervous system, cardiovascular and metabolic disorders. The
 CC oligomers are also used for detecting cell type differentiation. ABC00010
 CC -ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and ABI00010-ABI82073
 CC represent the oligomers described in the invention. NOTE: The sequence
 CC data for this patent did not form part of the printed specification, but
 CC was obtained in electronic format from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX

XX SQ Sequence 12 BP; 4 A; 0 C; 4 G; 4 T; 0 U; 0 Other;
 Query Match 61.2%; Score 10.4; DB 1; Length 12;
 Best Local Similarity 91.7%; Pred. No. 93;
 Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 6 CAACACTCCTAT 17
 ||||| |||||
 DB 12 CAACATTCCTAT 1

RESULT 43
 ABCS1750/c
 ID ABCS1750 standard; DNA; 13 BP.
 XX
 AC ABCS1750;
 XX
 XX 21-FEB-2002 (first entry)
 XX
 DE Oligonucleotide SEQ ID NO 51767 for detecting SNP TSC0014432.
 XX
 KW SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
 KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
 KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
 XX
 OS Homo sapiens.
 XX
 PN WO200177384-A2.
 XX
 PD 18-OCT-2001.
 XX
 PF 06-APR-2001; 2001WO-IB0000713.
 XX
 PR 07-APR-2000; 2000DE-01019173.
 XX
 XX (EPIG-) EPIGENOMICS AG.
 XX
 XX Olek A, Piepenbrock C, Berlin K;
 XX WPI; 2001-657177/75.
 XX
 XX Set of oligonucleotides, useful for diagnosis and cell typing, is
 PT designed to detect single-nucleotide polymorphisms and cytosine
 PT methylation status.
 XX
 XX Claim 1; SEQ ID NO 51767; 29pp + Sequence Listing; German.
 XX

XX SQ Sequence 13 BP; 4 A; 0 C; 4 G; 5 T; 0 U; 0 Other;
 Query Match 61.2%; Score 10.4; DB 1; Length 13;
 Best Local Similarity 91.7%; Pred. No. 1e+02;
 Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 6 CAACACTCCTAT 17
 ||||| |||||
 DB 13 CAACATTCCTAT 2

RESULT 44
 ABF59814/c
 ID ABF59814 standard; DNA; 13 BP.
 XX
 AC ABF59814;
 XX
 XX 21-FEB-2002 (first entry)
 XX
 DE Oligonucleotide SEQ ID NO 159811 for detecting SNP TSC0040225.
 XX
 KW SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
 KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
 KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
 XX
 OS Homo sapiens.
 XX
 PN WO200177384-A2.
 XX
 PD 18-OCT-2001.
 XX
 PF 06-APR-2001; 2001WO-IB0000713.
 XX
 PR 07-APR-2000; 2000DE-01019173.
 XX
 XX (EPIG-) EPIGENOMICS AG.
 XX
 XX Olek A, Piepenbrock C, Berlin K;
 XX WPI; 2001-657177/75.
 XX
 XX Set of oligonucleotides, useful for diagnosis and cell typing, is
 PT designed to detect single-nucleotide polymorphisms and cytosine
 PT methylation status.
 XX
 XX Claim 1; SEQ ID NO 159811; 29pp + Sequence Listing; German.
 XX

XX SQ This invention describes novel oligonucleotide primers or peptide nucleic
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
 CC and cytosine methylation status in chemically pretreated genomic DNA. The
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a
 CC range of diseases including immune system, gastrointestinal, respiratory,
 CC central nervous system, cardiovascular and metabolic disorders. The
 CC oligomers are also used for detecting cell type differentiation. ABC00010
 CC -ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and ABI00010-ABI82073
 CC represent the oligomers described in the invention. NOTE: The sequence
 CC data for this patent did not form part of the printed specification, but
 CC was obtained in electronic format from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX

XX SQ Claim 1; SEQ ID NO 159811; 29pp + Sequence Listing; German.
 XX
 XX This invention describes novel oligonucleotide primers or peptide nucleic
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
 CC and cytosine methylation status in chemically pretreated genomic DNA. The
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a
 CC range of diseases including immune system, gastrointestinal, respiratory,
 CC central nervous system, cardiovascular and metabolic disorders. The
 CC oligomers are also used for detecting cell type differentiation. ABC00010
 CC -ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and ABI00010-ABI82073
 CC represent the oligomers described in the invention. NOTE: The sequence
 CC data for this patent did not form part of the printed specification, but
 CC was obtained in electronic format from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX

```

SQ Sequence 13 BP; 1 A; 0 C; 6 G; 5 T; 0 U; 1 Other;
Query Match 61.2%; Score 10.4; DB 1; Length 13;
Best Local Similarity 91.7%; Pred. No. 1e+02;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 CAGCAACTCTCC 14
Db 12 CAACAACACTCC 1

RESULT 45
ABF91047
XX ABF91047 standard; DNA; 13 BP.
AC ABF91047;
XX
XX
XX 22-FEB-2002 (first entry)
DT
DE Oligonucleotide SEQ ID NO 191044 for detecting SNP TSC0047001.
XX
XX SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
XX
XX Homo sapiens.
OS
XX WO200177384-A2.
PN
XX 18-OCT-2001.
PD
XX
XX 22-FEB-2002 (first entry)
DT
DE Oligonucleotide SEQ ID NO 191044 for detecting SNP TSC0047001.
XX
XX SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
XX
XX Homo sapiens.
OS
XX WO200177384-A2.
PN
XX 18-OCT-2001.
PD
XX
XX 06-APR-2001; 2001WO-IB000713.
PF
XX 07-APR-2000; 2000DE-01019173.
PR
XX (EPIG-) EPIGENOMICS AG.
PA
PI Olek A, Piepenbrock C, Berlin K;
XX WPI; 2001-657177/75.
DR
XX
XX Set of oligonucleotides, useful for diagnosis and cell typing, is
PT designed to detect single-nucleotide polymorphisms and cytosine
PT methylation status.
XX
XX Claim 1; SEQ ID NO 191044; 29pp + Sequence Listing; German.
PS
XX This invention describes novel oligonucleotide primers or peptide nucleic
XX acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
XX and cytosine methylation status in chemically pretreated genomic DNA. The
XX oligonucleotides are used for diagnosis and/or prognosis of cancer and a
XX range of diseases including immune system, gastrointestinal, respiratory,
XX central nervous system, cardiovascular and metabolic disorders. The
XX oligomers are also used for detecting cell type differentiation. ABC00010
XX -ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and AB100010-AB182073
XX represent the oligomers described in the invention. NOTE: The sequence
XX data for this patent did not form part of the printed specification, but
XX was obtained in electronic format from WIPO at
XX ftp.wipo.int/pub/published_pct_sequences
XX
XX Sequence 13 BP; 3 A; 0 C; 6 G; 4 T; 0 U; 0 Other;
SQ
Query Match 61.2%; Score 10.4; DB 1; Length 13;
Best Local Similarity 91.7%; Pred. No. 1e+02;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 6 CAACACTCTCTAT 17
Db 2 CACCACTCTCTAT 13

RESULT 46
ABH23826/c
XX
XX
XX
XX 22-FEB-2002 (first entry)
DT
DE Oligonucleotide SEQ ID NO 163329 for detecting SNP TSC0041059.
XX
XX SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
XX
XX Homo sapiens.
OS

```


CC central nervous system, cardiovascular and metabolic disorders. The
 CC oligomers are also used for detecting cell type differentiation. ABC00010
 CC -ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and ABI00010-ABI82073
 CC represent the oligomers described in the invention. NOTE: The sequence
 CC data for this patent did not form part of the printed specification, but
 CC was obtained in electronic format from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences

XX SQ Sequence 13 BP; 5 A; 6 C; 0 G; 2 T; 0 U; 0 Other;

Query Match 61.2%; Score 10.4; DB 1; Length 13;
 Best Local Similarity 91.7%; Pred. No. 1e+02; Mismatches 0; Gaps 0;
 Matches 11; Conservative 0; Indels 1; Indels 0; Gaps 0;

Qy 4 AGCAACTCTCT 15
 Db 2 ACCAACAACCTCT 13

RESULT 50

ABH62283
 ID ABH62283 standard; DNA; 13 BP.

XX AC ABH62283;

DT 22-FEB-2002 (first entry)

XX DE Oligonucleotide SEQ ID NO 262260 for detecting SNP TSC0010773.

XX SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
 XX peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
 XX central nervous system; gastrointestinal; respiratory; immune; metabolic.

XX OS Homo sapiens.

XX PN WO200177384-A2.

XX PD 18-OCT-2001.

XX PF 06-APR-2001; 2001WO-IB000713.

XX PR 07-APR-2000; 2000DE-01019173.

XX PA (BFIG-) BFIGENOMICS AG.

XX PI Olek A, Piepenbrock C, Berlin K;

XX DR WPI; 2001-657177/75.

XX PT Set of oligonucleotides, useful for diagnosis and cell typing, is
 XX designed to detect single-nucleotide polymorphisms and cytosine
 XX methylation status.

XX PS Claim 1; SEQ ID NO 262260; 29pp + Sequence Listing; German.

XX This invention describes novel oligonucleotide primers or peptide nucleic
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
 CC and cytosine methylation status in chemically pretreated genomic DNA. The
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a
 CC range of diseases including immune system, gastrointestinal, respiratory,
 CC central nervous system, cardiovascular and metabolic disorders. The
 CC oligomers are also used for detecting cell type differentiation. ABC00010
 CC -ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and ABI00010-ABI82073
 CC represent the oligomers described in the invention. NOTE: The sequence
 CC data for this patent did not form part of the printed specification, but
 CC was obtained in electronic format from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences

XX SQ Sequence 13 BP; 3 A; 6 C; 0 G; 4 T; 0 U; 0 Other;

Query Match 61.2%; Score 10.4; DB 1; Length 13;
 Best Local Similarity 91.7%; Pred. No. 1e+02; Mismatches 1; Indels 0; Gaps 0;
 Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 6 CAACACTCTCTAT 17
 Db 1 CAACCTCTCTAT 12

Search completed: November 21, 2006, 14:29:06
 Job time : 0.001 secs

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OM nucleic - nucleic search, using sw model

Run on: November 21, 2006, 14:33:58 ; Search time 0.001 Seconds
(without alignments)
6.358 Million cell updates/sec

Title: US-10-823-197-7
Perfect score: 17
Sequence: 1 CTCAGCAACTCCTAT 17

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 0.5

Searched: 16 seqs, 187 residues

Total number of hits satisfying chosen parameters: 32

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 16 summaries

Database : rnpbndb.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	17	100.0	17	1	US-11-203-682-90
2	17	100.0	17	1	US-11-283-168-1
3	8.4	49.4	11	1	US-11-364-118-4
4	8.4	49.4	11	1	US-11-364-118-229
C 5	8.4	49.4	11	1	US-11-158-209-97
C 6	8.4	49.4	11	1	US-11-158-209-545
C 7	8.4	49.4	12	1	US-11-212-812A-395
C 8	8.4	49.4	12	1	US-11-212-386A-395
C 9	8	47.1	11	1	US-11-364-118-390
C 10	8	47.1	11	1	US-11-364-118-457
C 11	7.8	45.9	11	1	US-11-158-209-691
C 12	7.8	45.9	11	1	US-11-158-209-1221
C 13	7.8	45.9	11	1	US-11-148-303-66
C 14	7.4	43.5	10	1	US-10-691-012-35
C 15	7.4	43.5	10	1	US-11-148-303-23
C 16	7.4	43.5	10	1	US-11-148-303-128

ALIGNMENTS

RESULT 1
US-11-203-682-90
; Sequence 90, Application US/11203682
; Publication No. US20060183161A1
; GENERAL INFORMATION:
; APPLICANT: Nicklin, Martin
; APPLICANT: Barton, Jenny
; TITLE OF INVENTION: IL-11 GENE AND POLYPEPTIDE PRODUCTS
; FILE REFERENCE: 24299-517 CIP
; CURRENT APPLICATION NUMBER: US/11/203,682
; CURRENT FILING DATE: 2005-08-12
; PRIOR APPLICATION NUMBER: 09/617,720

; PRIOR FILING DATE: 2000-07-17
; NUMBER OF SEQ ID NOS: 132
; SOFTWARE: PatentIn Ver. 3.2
; SEQ ID NO 90
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Chemically synthesized primer
US-11-203-682-90

Query Match 100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACTCCTAT 17
| | | | | | | | | | | | | | | | | |
Db 1 CTCAGCAACTCCTAT 17

RESULT 2

US-11-283-168-1
; Sequence 1, Application US/11283168
; Publication No. US20060252055A1
; GENERAL INFORMATION:
; APPLICANT: Francis, Sheila E
; APPLICANT: Crossman, David C
; APPLICANT: Duff, Gordon W
; APPLICANT: Kornman, Kenneth S
; APPLICANT: Martinez, Katherine
; TITLE OF INVENTION: DIAGNOSTICS AND THERAPEUTICS FOR CARDIOVASCULAR DISORDERS
; FILE REFERENCE: 24299-504 CIP2 CON CIP1
; CURRENT APPLICATION NUMBER: US/11/283,168
; CURRENT FILING DATE: 2005-11-17
; PRIOR APPLICATION NUMBER: 10/320,360
; PRIOR FILING DATE: 2002-12-13
; PRIOR APPLICATION NUMBER: 09/431,352
; PRIOR FILING DATE: 1999-11-01
; PRIOR APPLICATION NUMBER: 09/320,395
; PRIOR FILING DATE: 1999-05-26
; PRIOR APPLICATION NUMBER: 08/813,456
; PRIOR FILING DATE: 1997-03-10
; NUMBER OF SEQ ID NOS: 18
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 1
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Artificial
; FEATURE:
; OTHER INFORMATION: Primer
US-11-283-168-1

Query Match 100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACTCCTAT 17
| | | | | | | | | | | | | | | | | |
Db 1 CTCAGCAACTCCTAT 17

RESULT 3

US-11-364-118-4
; Sequence 4, Application US/11364118
; Publication No. US20060204992A1
; GENERAL INFORMATION:
; APPLICANT: Olaf Holtkotter
; APPLICANT: Dirk Petersohn
; APPLICANT: Kordula Schlotmann
; APPLICANT: Melanie Giesen
; APPLICANT: Daniela Kessler-Becker
; TITLE OF INVENTION: Method for Determining Hair Cycle Markers
; FILE REFERENCE: H 06059 PCT

```

; CURRENT APPLICATION NUMBER: US/11/364.118
; CURRENT FILING DATE: 2006-02-28
; PRIOR APPLICATION NUMBER: PCT/EP2004/009435
; PRIOR FILING DATE: 2004-08-24
; PRIOR APPLICATION NUMBER: 103 40 373.6-41
; PRIOR FILING DATE: 2003-08-30
; NUMBER OF SEQ ID NOS: 570
; SOFTWARE: SeqWin99, version 1.02
; SEQ ID NO 4
; LENGTH: 11
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-364-118-4

Query Match      49.4%; Score 8.4; DB 1; Length 11;
Best Local Similarity 90.0%; Pred. No. 6.6;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      6 CAACACTCCT 15
DB      1 CAACATTCCT 10

RESULT 4
US-11-364-118-229
; Sequence 229, Application US/11364118
; Publication No. US20060204992A1
; GENERAL INFORMATION:
; APPLICANT: Olaf Holtkotter
; APPLICANT: Dirk Petersohn
; APPLICANT: Kordula Schlottmann
; APPLICANT: Melanie Kessler-Becker
; TITLE OF INVENTION: Method for Determining Hair Cycle Markers
; FILE REFERENCE: H 06059 PCT
; CURRENT APPLICATION NUMBER: US/11/364.118
; CURRENT FILING DATE: 2006-02-28
; PRIOR APPLICATION NUMBER: PCT/EP2004/009435
; PRIOR FILING DATE: 2004-08-24
; PRIOR APPLICATION NUMBER: 103 40 373.6-41
; PRIOR FILING DATE: 2003-08-30
; NUMBER OF SEQ ID NOS: 570
; SOFTWARE: SeqWin99, version 1.02
; SEQ ID NO 229
; LENGTH: 11
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-364-118-229

Query Match      49.4%; Score 8.4; DB 1; Length 11;
Best Local Similarity 90.0%; Pred. No. 6.6;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      6 CAACACTCCT 15
DB      1 CAACATTCCT 10

RESULT 5
US-11-158-209-97/c
; Sequence 97, Application US/11158209
; Publication No. US2006008852A1
; GENERAL INFORMATION:
; APPLICANT: Dirk Petersohn
; APPLICANT: Kordula Schlottmann
; APPLICANT: Thomas Gassenmeier
; APPLICANT: Olaf Holtkotter
; APPLICANT: Marcus Conradt
; APPLICANT: Kay Hofmann
; TITLE OF INVENTION: Method for Determining the Homeostasis of Hairy Skin
; FILE REFERENCE: H 05667 PCT
; CURRENT APPLICATION NUMBER: US/11/158.209
; CURRENT FILING DATE: 2005-06-20

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; PRIOR APPLICATION NUMBER: PCT/EP2003/014070
; PRIOR FILING DATE: 2003-12-11
; PRIOR APPLICATION NUMBER: 102 60 931.4-41
; PRIOR FILING DATE: 2002-12-20
; NUMBER OF SEQ ID NOS: 1335
; SOFTWARE: SeqWin99, version 1.02
; SEQ ID NO 97
; LENGTH: 11
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-158-209-97

Query Match      49.4%; Score 8.4; DB 1; Length 11;
Best Local Similarity 90.0%; Pred. No. 6.6;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1 CTCAGCAACA 10
DB      11 CCAGCAACA 2

RESULT 6
US-11-158-209-545/c
; Sequence 545, Application US/11158209
; Publication No. US2006008852A1
; GENERAL INFORMATION:
; APPLICANT: Dirk Petersohn
; APPLICANT: Kordula Schlottmann
; APPLICANT: Thomas Gassenmeier
; APPLICANT: Olaf Holtkotter
; APPLICANT: Marcus Conradt
; APPLICANT: Kay Hofmann
; TITLE OF INVENTION: Method for Determining the Homeostasis of Hairy Skin
; FILE REFERENCE: H 05667 PCT
; CURRENT APPLICATION NUMBER: US/11/158.209
; CURRENT FILING DATE: 2005-06-20
; PRIOR APPLICATION NUMBER: PCT/EP2003/014070
; PRIOR FILING DATE: 2003-12-11
; PRIOR APPLICATION NUMBER: 102 60 931.4-41
; PRIOR FILING DATE: 2002-12-20
; NUMBER OF SEQ ID NOS: 1335
; SOFTWARE: SeqWin99, version 1.02
; SEQ ID NO 545
; LENGTH: 11
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-158-209-545

Query Match      49.4%; Score 8.4; DB 1; Length 11;
Best Local Similarity 90.0%; Pred. No. 6.6;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      3 CAGCAACACT 12
DB      10 CAGCAAAACT 1

RESULT 7
US-11-212-812A-395
; Sequence 395, Application US/11212812A
; Publication No. US20060121452A1
; GENERAL INFORMATION:
; APPLICANT: Dhallan, Ravinder S.
; TITLE OF INVENTION: METHODS FOR DETECTION OF GENETIC
; FILE REFERENCE: DISORDERS
; FILE REFERENCE: 543312000401
; CURRENT APPLICATION NUMBER: US/11/212.812A
; CURRENT FILING DATE: 2005-08-26
; PRIOR APPLICATION NUMBER: PCT/US04/006337
; PRIOR FILING DATE: 2004-03-01
; PRIOR APPLICATION NUMBER: PCT/US03/06198
; PRIOR FILING DATE: 2003-02-28
; PRIOR APPLICATION NUMBER: US60/378,354

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; PRIOR FILING DATE: 2002-05-08
; NUMBER OF SEQ ID NOS: 725
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 395
; LENGTH: 12
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Primer
US-11-212-812A-395

Query Match 49.4%; Score 8.4; DB 1; Length 12;
Best Local Similarity 90.0%; Pred. No. 6.1;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 TCAGCAACAC 11
Db 1 TCAGTAACAC 10

RESULT 8
US-11-212-386A-395
; Sequence 395, Application US/11212386A
; Publication No. US20060160105A1
; GENERAL INFORMATION:
; APPLICANT: Dhallan, Ravinder S.
; TITLE OF INVENTION: METHODS FOR DETECTION OF GENETIC
; FILE REFERENCE: 54331200402
; CURRENT APPLICATION NUMBER: US/11/212,386A
; CURRENT FILING DATE: 2005-08-26
; PRIOR APPLICATION NUMBER: US10/661,165
; PRIOR FILING DATE: 2003-09-11
; PRIOR APPLICATION NUMBER: PCT/US03/27308
; PRIOR FILING DATE: 2003-08-29
; PRIOR APPLICATION NUMBER: PCT/US03/06198
; PRIOR FILING DATE: 2003-02-28
; PRIOR APPLICATION NUMBER: US60/378,354
; PRIOR FILING DATE: 2002-08-08
; NUMBER OF SEQ ID NOS: 725
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 395
; LENGTH: 12
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Primer
US-11-212-386A-395

Query Match 49.4%; Score 8.4; DB 1; Length 12;
Best Local Similarity 90.0%; Pred. No. 6.1;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 TCAGCAACAC 11
Db 1 TCAGTAACAC 10

RESULT 9
US-11-364-118-390/c
; Sequence 390, Application US/11364118
; Publication No. US20060204992A1
; GENERAL INFORMATION:
; APPLICANT: Olaf Holtkötter
; APPLICANT: Dirk Petersohn
; APPLICANT: Kordula Schlotmann
; APPLICANT: Daniela Kessler-Becker
; TITLE OF INVENTION: Method for Determining Hair Cycle Markers
; FILE REFERENCE: H 06059 PCT
; CURRENT APPLICATION NUMBER: US/11/364,118
; CURRENT FILING DATE: 2006-02-28
; PRIOR APPLICATION NUMBER: PCT/EP2004/009435

; PRIOR FILING DATE: 2004-08-24
; PRIOR APPLICATION NUMBER: 103 40 373.6-41
; PRIOR FILING DATE: 2003-08-30
; NUMBER OF SEQ ID NOS: 570
; SOFTWARE: SeqWin99, version 1.02
; SEQ ID NO 390
; LENGTH: 11
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-364-118-390

Query Match 47.1%; Score 8; DB 1; Length 11;
Best Local Similarity 100.0%; Pred. No. 7.5;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 8 ACACCTCCT 15
Db 9 ACACCTCCT 2

RESULT 10
US-11-364-118-457/c
; Sequence 457, Application US/11364118
; Publication No. US20060204992A1
; GENERAL INFORMATION:
; APPLICANT: Olaf Holtkötter
; APPLICANT: Dirk Petersohn
; APPLICANT: Kordula Schlotmann
; APPLICANT: Daniela Kessler-Becker
; TITLE OF INVENTION: Method for Determining Hair Cycle Markers
; FILE REFERENCE: H 06059 PCT
; CURRENT APPLICATION NUMBER: US/11/364,118
; CURRENT FILING DATE: 2006-02-28
; PRIOR APPLICATION NUMBER: PCT/EP2004/009435
; PRIOR FILING DATE: 2004-08-24
; PRIOR APPLICATION NUMBER: 103 40 373.6-41
; PRIOR FILING DATE: 2003-08-30
; NUMBER OF SEQ ID NOS: 570
; SOFTWARE: SeqWin99, version 1.02
; SEQ ID NO 457
; LENGTH: 11
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-364-118-457

Query Match 47.1%; Score 8; DB 1; Length 11;
Best Local Similarity 100.0%; Pred. No. 7.5;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 8 ACACCTCCT 15
Db 9 ACACCTCCT 2

RESULT 11
US-11-158-209-691/c
; Sequence 691, Application US/11158209
; Publication No. US2006008852A1
; GENERAL INFORMATION:
; APPLICANT: Dirk Petersohn
; APPLICANT: Kordula Schlotmann
; APPLICANT: Thomas Gassenmeier
; APPLICANT: Olaf Holtkötter
; APPLICANT: Marcus Conradt
; APPLICANT: Kay Hofmann
; TITLE OF INVENTION: Method for Determining the Homeostasis of Hairy Skin
; FILE REFERENCE: H 05667 PCT
; CURRENT APPLICATION NUMBER: US/11/158,209
; CURRENT FILING DATE: 2005-06-20
; PRIOR APPLICATION NUMBER: PCT/EP2003/014070
; PRIOR FILING DATE: 2003-12-11
; PRIOR APPLICATION NUMBER: 102 60 931.4-41

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; PRIOR FILING DATE: 2002-12-20
; NUMBER OF SEQ ID NOS: 1335
; SOFTWARE: SeqWin99, version 1.02
; SEQ ID NO 691
; LENGTH: 11
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-158-209-691

Query Match      45.9%; Score 7.8; DB 1; Length 11;
Best Local Similarity 81.8%; Pred. No. 7.9;
Matches 9; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy      4 AGCAACTCTCC 14
Db      11 AGCTGCATCTC 1

RESULT 12
US-11-158-209-1221/c
; Sequence 1221, Application US/11158209
; Publication No. US2006008852A1
; GENERAL INFORMATION:
; APPLICANT: Dirk Petersohn
; APPLICANT: Kordula Schlotmann
; APPLICANT: Thomas Gassenmeier
; APPLICANT: Olaf Holtkotter
; APPLICANT: Marcus Conrad
; APPLICANT: Kay Hofmann
; TITLE OF INVENTION: Method for Determining the Homeostasis of Hairy Skin
; FILE REFERENCE: H 05667 PCT
; CURRENT APPLICATION NUMBER: US/11/158,209
; CURRENT FILING DATE: 2005-06-20
; PRIOR APPLICATION NUMBER: PCT/EP2003/014070
; PRIOR FILING DATE: 2003-12-11
; PRIOR APPLICATION NUMBER: 102 60 931.4-41
; PRIOR FILING DATE: 2002-12-20
; NUMBER OF SEQ ID NOS: 1335
; SOFTWARE: SeqWin99, version 1.02
; SEQ ID NO 1221
; LENGTH: 11
; TYPE: DNA
; ORGANISM: Homo Sapiens
US-11-158-209-1221

Query Match      45.9%; Score 7.8; DB 1; Length 11;
Best Local Similarity 81.8%; Pred. No. 7.9;
Matches 9; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy      1 CTCAGCAACAC 11
Db      11 CTCAACTACAC 1

RESULT 13
US-11-148-303-66
; Sequence 66, Application US/11148303
; Publication No. US20060154886A1
; GENERAL INFORMATION:
; APPLICANT: Gruenthal GmbH
; TITLE OF INVENTION: Regulatory elements in the 5' region of the VR1 gene
; FILE REFERENCE: GR01P003WO
; CURRENT APPLICATION NUMBER: US/11/148,303
; CURRENT FILING DATE: 2005-06-09
; NUMBER OF SEQ ID NOS: 781
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 66
; LENGTH: 11
; TYPE: DNA
; ORGANISM: Rattus norvegicus
; FEATURE:
; OTHER INFORMATION: V$NFY Q6
US-11-148-303-66

Query Match      45.9%; Score 7.8; DB 1; Length 11;
Best Local Similarity 81.8%; Pred. No. 7.9;
Matches 9; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy      1 CTCAGCAACAC 11
Db      11 CTCAACTACAC 1

RESULT 14
US-10-691-012-35/c
; Sequence 35, Application US/10691012
; Publication No. US20060160731A1
; GENERAL INFORMATION:
; APPLICANT: Buchardt, Ole
; APPLICANT: Egholm, Michael
; APPLICANT: Nielsen, Peter Eigil
; APPLICANT: Berg, Rolf Henrik
; TITLE OF INVENTION: Peptide Nucleic Acids
; FILE REFERENCE: IS180540
; CURRENT APPLICATION NUMBER: US/10/691,012
; CURRENT FILING DATE: 2003-10-22
; PRIOR APPLICATION NUMBER: US/08/108,591
; PRIOR FILING DATE: 1993-11-22
; NUMBER OF SEQ ID NOS: 43
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 35
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Novel Sequence
US-10-691-012-35

Query Match      43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 9.8;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      9 CACTCCTAT 17
Db      10 CACTACTAT 2

RESULT 15
US-11-148-303-23
; Sequence 23, Application US/11148303
; Publication No. US20060154886A1
; GENERAL INFORMATION:
; APPLICANT: Gruenthal GmbH
; TITLE OF INVENTION: Regulatory elements in the 5' region of the VR1 gene
; FILE REFERENCE: GR01P003WO
; CURRENT APPLICATION NUMBER: US/11/148,303
; CURRENT FILING DATE: 2005-06-09
; NUMBER OF SEQ ID NOS: 781
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 23
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Rattus norvegicus
; FEATURE:
; OTHER INFORMATION: V$SOK5 01
US-11-148-303-23

Query Match      43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 9.8;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      6 CAACACTCC 14
Db      2 CAACAATCC 10
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RESULT 16
US-11-148-303-128
; Sequence 128, Application US/11148303
; Publication No. US20060154886A1
; GENERAL INFORMATION:
; APPLICANT: Gruenthal GmbH
; TITLE OF INVENTION: Regulatory elements in the 5' region of the VR1 gene
; FILE REFERENCE: GR01P003WO
; CURRENT APPLICATION NUMBER: US/11/148,303
; CURRENT FILING DATE: 2005-06-09
; NUMBER OF SEQ ID NOS: 781
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 128
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Mus musculus
; FEATURE:
; OTHER INFORMATION: VS0X5 01
US-11-148-303-128

Query Match      43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 9.8;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 5 GCACACTC 13
   |||||
Db 1 GCNACATC 9

Search completed: November 21, 2006, 14:33:58
Job time : 0.001 secs

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OM nucleic - nucleic search, using sw model

Run on: November 21, 2006, 14:32:49 ; Search time 0.001 Seconds
(without alignments)
117.640 Million cell updates/sec

Title: US-10-823-197-7

Perfect score: 17

Sequence: 1 CTCAGCAACTCTCTAT 17

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 0.5

Searched: 262 seqs, 3460 residues

Total number of hits satisfying chosen parameters: 524

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 262 summaries

Database : rnpbmdb.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	17	100.0	17	1	US-09-845-129-22 Sequence 22, Appl
2	17	100.0	17	1	US-09-888-056A-3 Sequence 3, Appl
3	17	100.0	17	1	US-10-167-127-23 Sequence 23, Appl
4	17	100.0	17	1	US-10-172-919-7 Sequence 7, Appl
5	17	100.0	17	1	US-10-320-360-1 Sequence 1, Appl
6	17	100.0	17	1	US-10-802-061-22 Sequence 22, Appl
7	17	100.0	17	1	US-10-823-197-7 Sequence 7, Appl
8	17	100.0	17	1	US-10-712-882-5 Sequence 5, Appl
9	17	100.0	17	1	US-10-838-503-9 Sequence 9, Appl
10	17	100.0	17	1	US-11-121-634-22 Sequence 22, Appl
11	14	82.4	17	1	US-10-626-830-9 Sequence 9, Appl
12	14	82.4	19	1	US-11-083-784-1215680 Sequence 1215680,
13	14	82.4	19	1	US-11-083-784-1215694 Sequence 1215694,
14	14	82.4	19	1	US-11-083-784-1215717 Sequence 1215717,
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21	13.8	81.2	19	1	US-11-101-244-921461 Sequence 921461,
22	13.8	81.2	19	1	US-11-101-244-1372880 Sequence 1372880,
23	13.4	78.8	19	1	US-11-083-784-628637 Sequence 628637,
24	13.4	78.8	19	1	US-11-083-784-628845 Sequence 628845,
25	13.4	78.8	19	1	US-11-101-244-628637 Sequence 628637,
26	13.4	78.8	19	1	US-11-101-244-628845 Sequence 628845,
27	12.8	75.3	18	1	US-10-310-914A-273305 Sequence 273305,
28	12.4	72.9	16	1	US-10-096-125-14 Sequence 14, Appl
29	11	64.7	12	1	US-10-257-017B-302368 Sequence 302368,
30	11	64.7	13	1	US-10-257-017B-71093 Sequence 71093, A
31	11	64.7	13	1	US-10-257-017B-71094 Sequence 71094, A
32	11	64.7	13	1	US-10-257-017B-136017 Sequence 136017,
33	11	64.7	13	1	US-10-257-017B-136018 Sequence 136018,

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US-10-257-017B-315837

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Sequence 282853,
Sequence 283403,
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Sequence 315837,

107	9.4	55.3	12	1	US-10-257-017B-321267	Sequence 321267,	C 180	9.4	55.3	13	1	US-10-257-017B-178081	Sequence 178081,
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C 110	9.4	55.3	12	1	US-10-257-017B-32452	Sequence 32452,	C 183	9.4	55.3	13	1	US-10-257-017B-179042	Sequence 179042,
C 111	9.4	55.3	12	1	US-10-257-017B-337997	Sequence 337997,	C 184	9.4	55.3	13	1	US-10-257-017B-190349	Sequence 190349,
C 112	9.4	55.3	12	1	US-10-257-017B-351387	Sequence 351387,	C 185	9.4	55.3	13	1	US-10-257-017B-190350	Sequence 190350,
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C 117	9.4	55.3	12	1	US-10-257-017B-379316	Sequence 379316,	C 190	9.4	55.3	13	1	US-10-257-017B-211433	Sequence 211433,
C 118	9.4	55.3	13	1	US-10-257-017B-309	Sequence 309, App	C 191	9.4	55.3	13	1	US-10-257-017B-211434	Sequence 211434,
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C 120	9.4	55.3	13	1	US-10-257-017B-24595	Sequence 24595, A	C 193	9.4	55.3	13	1	US-10-257-017B-226550	Sequence 226550,
C 121	9.4	55.3	13	1	US-10-257-017B-24596	Sequence 24596, A	C 194	9.4	55.3	13	1	US-10-257-017B-227085	Sequence 227085,
C 122	9.4	55.3	13	1	US-10-257-017B-25817	Sequence 25817, A	C 195	9.4	55.3	13	1	US-10-257-017B-227086	Sequence 227086,
C 123	9.4	55.3	13	1	US-10-257-017B-25818	Sequence 25818, A	C 196	9.4	55.3	13	1	US-10-257-017B-227087	Sequence 227087,
C 124	9.4	55.3	13	1	US-10-257-017B-25818	Sequence 25818, A	C 197	9.4	55.3	13	1	US-10-257-017B-227088	Sequence 227088,
C 125	9.4	55.3	13	1	US-10-257-017B-39645	Sequence 39645, A	C 198	9.4	55.3	13	1	US-10-257-017B-227088	Sequence 227088,
C 126	9.4	55.3	13	1	US-10-257-017B-39646	Sequence 39646, A	C 199	9.4	55.3	13	1	US-10-257-017B-231119	Sequence 231119,
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C 131	9.4	55.3	13	1	US-10-257-017B-50897	Sequence 50897, A	C 204	9.4	55.3	13	1	US-10-257-017B-235004	Sequence 235004,
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C 140	9.4	55.3	13	1	US-10-257-017B-71102	Sequence 71102, A	C 213	9.4	55.3	13	1	US-10-257-017B-274536	Sequence 274536,
C 141	9.4	55.3	13	1	US-10-257-017B-75403	Sequence 75403, A	C 214	9.4	55.3	13	1	US-10-257-017B-289056	Sequence 289056,
C 142	9.4	55.3	13	1	US-10-257-017B-75404	Sequence 75404, A	C 215	9.4	55.3	13	1	US-10-257-017B-304514	Sequence 304514,
C 143	9.4	55.3	13	1	US-10-257-017B-78917	Sequence 78917, A	C 216	9.4	55.3	13	1	US-10-257-017B-304514	Sequence 304514,
C 144	9.4	55.3	13	1	US-10-257-017B-78918	Sequence 78918, A	C 217	9.4	55.3	13	1	US-10-257-017B-308500	Sequence 308500,
C 145	9.4	55.3	13	1	US-10-257-017B-86482	Sequence 86482, A	C 218	9.4	55.3	13	1	US-10-257-017B-318085	Sequence 318085,
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C 149	9.4	55.3	13	1	US-10-257-017B-96645	Sequence 96645, A	C 222	9.4	55.3	13	1	US-10-257-017B-349259	Sequence 349259,
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C 153	9.4	55.3	13	1	US-10-257-017B-108233	Sequence 108233,	C 226	9.4	55.3	13	1	US-10-257-017B-271995	Sequence 271995,
C 154	9.4	55.3	13	1	US-10-257-017B-108234	Sequence 108234,	C 227	9.4	55.3	13	1	US-10-257-017B-277739	Sequence 277739,
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C 157	9.4	55.3	13	1	US-10-257-017B-117651	Sequence 117651,	C 230	9.4	55.3	13	1	US-10-257-017B-290054	Sequence 290054,
C 158	9.4	55.3	13	1	US-10-257-017B-117652	Sequence 117652,	C 231	9.4	55.3	13	1	US-10-257-017B-290876	Sequence 290876,
C 159	9.4	55.3	13	1	US-10-257-017B-127381	Sequence 127381,	C 232	9.4	55.3	13	1	US-10-257-017B-295767	Sequence 295767,
C 160	9.4	55.3	13	1	US-10-257-017B-127382	Sequence 127382,	C 233	9.4	55.3	13	1	US-10-257-017B-298432	Sequence 298432,
C 161	9.4	55.3	13	1	US-10-257-017B-127385	Sequence 127385,	C 234	9.4	55.3	13	1	US-10-257-017B-303022	Sequence 303022,
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C 163	9.4	55.3	13	1	US-10-257-017B-132701	Sequence 132701,	C 236	9.4	55.3	13	1	US-10-257-017B-307496	Sequence 307496,
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C 165	9.4	55.3	13	1	US-10-257-017B-136015	Sequence 136015,	C 238	9.4	55.3	13	1	US-10-257-017B-314086	Sequence 314086,
C 166	9.4	55.3	13	1	US-10-257-017B-136016	Sequence 136016,	C 239	9.4	55.3	13	1	US-10-257-017B-314087	Sequence 314087,
C 167	9.4	55.3	13	1	US-10-257-017B-153289	Sequence 153289,	C 240	9.4	55.3	13	1	US-10-257-017B-321092	Sequence 321092,
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C 169	9.4	55.3	13	1	US-10-257-017B-153995	Sequence 153995,	C 242	9.4	55.3	13	1	US-10-257-017B-322014	Sequence 322014,
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C 171	9.4	55.3	13	1	US-10-257-017B-157065	Sequence 157065,	C 244	9.4	55.3	13	1	US-10-257-017B-326438	Sequence 326438,
C 172	9.4	55.3	13	1	US-10-257-017B-157066	Sequence 157066,	C 245	9.4	55.3	13	1	US-10-257-017B-328173	Sequence 328173,
C 173	9.4	55.3	13	1	US-10-257-017B-169365	Sequence 169365,	C 246	9.4	55.3	13	1	US-10-257-017B-329594	Sequence 329594,
C 174	9.4	55.3	13	1	US-10-257-017B-169366	Sequence 169366,	C 247	9.4	55.3	13	1	US-10-257-017B-332773	Sequence 332773,
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C 178	9.4	55.3	13	1	US-10-257-017B-174910	Sequence 174910,	C 251	9.4	55.3	13	1	US-10-257-017B-352697	Sequence 352697,
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c 253 8.8 51.8 12 1 US-10-257-017B-360500 Sequence 360500,
 c 254 8.8 51.8 12 1 US-10-257-017B-362463 Sequence 362463,
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 c 258 8.8 51.8 12 1 US-10-257-017B-371753 Sequence 371753,
 c 259 8.8 51.8 12 1 US-10-257-017B-374492 Sequence 374492,
 c 260 8.8 51.8 12 1 US-10-257-017B-375813 Sequence 375813,
 c 261 8.8 51.8 12 1 US-10-257-017B-379818 Sequence 379818,
 c 262 8.8 51.8 12 1 US-10-257-017B-379853 Sequence 379853,

ALIGNMENTS

RESULT 1
 US-09-845-129-22
 ; Sequence 22, Application US/09845129
 ; Patent No. US20020146700A1
 ; GENERAL INFORMATION:
 ; APPLICANT: DUFF, GORDON W.
 ; APPLICANT: COX, ANGELA
 ; APPLICANT: CAMP, NICOLA J.
 ; TITLE OF INVENTION: DIGTOVINE, FRANCESCO S.
 ; TITLE OF INVENTION: DIAGNOSTICS AND THERAPEUTICS FOR DISEASES ASSOCIATED
 ; WITH AN IL-1 INFLAMMATORY HAPLOTYPE
 ; FILE REFERENCE: MSA-010.02
 ; CURRENT APPLICATION NUMBER: US/09/845,129
 ; CURRENT FILING DATE: 2001-04-27
 ; PRIOR APPLICATION NUMBER: 09/345,217
 ; PRIOR FILING DATE: 1999-06-30
 ; PRIOR APPLICATION NUMBER: PCT/GB98/01481
 ; PRIOR FILING DATE: 1998-05-21
 ; PRIOR APPLICATION NUMBER: 9711040.7
 ; PRIOR FILING DATE: 1997-05-29
 ; NUMBER OF SEQ ID NOS: 32
 ; SOFTWARE: PatentIn Ver. 2.0
 ; SEQ ID NO 22
 ; LENGTH: 17
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: Description of Artificial Sequence: primer
 US-09-845-129-22

Query Match 100.0%; Score 17; DB 1; Length 17;
 Best Local Similarity 100.0%; Pred. No. 14;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGCAACACTCCTAT 17
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 DB 1 CTCAGCAACACTCCTAT 17

RESULT 2
 US-09-888-056A-3
 ; Sequence 3, Application US/09888056A
 ; Publication No. US20030124524A1
 ; GENERAL INFORMATION:
 ; APPLICANT: KORNMAN, KENNETH S.
 ; APPLICANT: DUFF, GORDON W.
 ; TITLE OF INVENTION: SCREENING ASSAYS FOR IDENTIFYING MODULATORS OF THE
 ; FILE REFERENCE: MSA-023.01
 ; CURRENT APPLICATION NUMBER: US/09/888,056A
 ; CURRENT FILING DATE: 2002-05-06
 ; PRIOR APPLICATION NUMBER: 60/213,853
 ; PRIOR FILING DATE: 2000-06-23
 ; NUMBER OF SEQ ID NOS: 30
 ; SOFTWARE: PatentIn Ver. 2.1
 ; SEQ ID NO 3
 ; LENGTH: 17
 ; TYPE: DNA

; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: Description of Artificial Sequence: Primer
 US-09-888-056A-3

Query Match 100.0%; Score 17; DB 1; Length 17;
 Best Local Similarity 100.0%; Pred. No. 14;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGCAACACTCCTAT 17
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 DB 1 CTCAGCAACACTCCTAT 17

RESULT 3
 US-10-167-127-23
 ; Sequence 23, Application US/10167127
 ; Publication No. US20030100031A1
 ; GENERAL INFORMATION:
 ; APPLICANT: DOWER, STEVEN
 ; APPLICANT: DUFF, GORDON W.
 ; TITLE OF INVENTION: INTEGRATIVE ASSAYS FOR MONITORING MOLECULAR ASSEMBLY
 ; TITLE OF INVENTION: EVENTS
 ; FILE REFERENCE: MSA-026.01 (20974-3601)
 ; CURRENT APPLICATION NUMBER: US/10/167,127
 ; CURRENT FILING DATE: 2002-06-11
 ; PRIOR APPLICATION NUMBER: 60/297,305
 ; PRIOR FILING DATE: 2001-06-11
 ; NUMBER OF SEQ ID NOS: 32
 ; SOFTWARE: PatentIn Ver. 2.1
 ; SEQ ID NO 23
 ; LENGTH: 17
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: Description of Artificial Sequence: Primer
 US-10-167-127-23

Query Match 100.0%; Score 17; DB 1; Length 17;
 Best Local Similarity 100.0%; Pred. No. 14;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGCAACACTCCTAT 17
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 DB 1 CTCAGCAACACTCCTAT 17

RESULT 4
 US-10-172-919-7
 ; Sequence 7, Application US/10172919
 ; Publication No. US20030152947A1
 ; GENERAL INFORMATION:
 ; APPLICANT: CROSSMAN, DAVID C.
 ; APPLICANT: DUFF, GORDON W.
 ; APPLICANT: FRANCIS, SHEILA E.
 ; APPLICANT: KORNMAN, KENNETH S.
 ; APPLICANT: BARNETT, KATHERINE
 ; TITLE OF INVENTION: METHODS FOR DETECTING AND TREATING THE EARLY ONSET OF
 ; FILE REFERENCE: MSA-025.01
 ; CURRENT APPLICATION NUMBER: US/10/172,919
 ; CURRENT FILING DATE: 2002-06-17
 ; PRIOR APPLICATION NUMBER: 60/298,493
 ; PRIOR FILING DATE: 2001-06-15
 ; NUMBER OF SEQ ID NOS: 31
 ; SOFTWARE: PatentIn Ver. 2.1
 ; SEQ ID NO 7
 ; LENGTH: 17
 ; TYPE: DNA
 ; ORGANISM: Artificial Sequence
 ; FEATURE:
 ; OTHER INFORMATION: Description of Artificial Sequence: Primer
 US-10-172-919-7

Query Match 100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 14;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
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Db 1 CTCAGCAACACTCCTAT 17

RESULT 5
US-10-320-360-1
; Sequence 1, Application US/10320360
; Publication No. US20030175764A1
; GENERAL INFORMATION:
; APPLICANT: Francis, Sheila E.
; APPLICANT: Crossman, David C.
; APPLICANT: Duff, Gordon W.
; APPLICANT: Kornman, Kenneth S.
; APPLICANT: Stephenson, Katherine
; TITLE OF INVENTION: DIAGNOSTICS AND THERAPEUTICS FOR CARDIOVASCULAR
; TITLE OF INVENTION: DISORDERS
; FILE REFERENCE: MSA-006.03
; CURRENT APPLICATION NUMBER: US/10/320,360
; CURRENT FILING DATE: 2002-12-13
; PRIOR APPLICATION NUMBER: 09/320,395
; PRIOR FILING DATE: 1999-05-26
; NUMBER OF SEQ ID NOS: 18
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: primer
US-10-320-360-1

Query Match 100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 14;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
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Db 1 CTCAGCAACACTCCTAT 17

RESULT 6
US-10-802-061-22
; Sequence 22, Application US/10802061
; Publication No. US20040152124A1
; GENERAL INFORMATION:
; APPLICANT: DUFF, GORDON W.
; APPLICANT: COX, ANGELA
; APPLICANT: CAMP, NICOLA J.
; APPLICANT: DIGIOVINE, FRANCESCO S.
; TITLE OF INVENTION: DIAGNOSTICS AND THERAPEUTICS FOR DISEASES ASSOCIATED
; TITLE OF INVENTION: WITH AN IL-1 INFLAMMATORY HAPLOTYPE
; FILE REFERENCE: 24299-508CON3
; CURRENT APPLICATION NUMBER: US/10/802,061
; CURRENT FILING DATE: 2004-03-15
; PRIOR APPLICATION NUMBER: 09/845,129
; PRIOR FILING DATE: 2001-04-27
; PRIOR APPLICATION NUMBER: 09/345,217
; PRIOR FILING DATE: 1999-06-30
; PRIOR APPLICATION NUMBER: PCT/GB98/01481
; PRIOR FILING DATE: 1998-05-21
; PRIOR APPLICATION NUMBER: 9711040.7
; PRIOR FILING DATE: 1997-05-29
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 22
; LENGTH: 17
; TYPE: DNA

; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: primer
US-10-802-061-22

Query Match 100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 14;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
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Db 1 CTCAGCAACACTCCTAT 17

RESULT 7
US-10-823-197-7
; Sequence 7, Application US/10823197
; Publication No. US20040229264A1
; GENERAL INFORMATION:
; APPLICANT: CROSSMAN, DAVID C.
; APPLICANT: DUFF, GORDON W.
; APPLICANT: FRANCIS, SHEILA E.
; APPLICANT: KORNMAN, KENNETH S.
; APPLICANT: STEPHENSON, KATHERINE
; TITLE OF INVENTION: DIAGNOSTICS AND THERAPEUTICS FOR RESTENOSIS
; FILE REFERENCE: 24299-514CIP2A DIV
; CURRENT APPLICATION NUMBER: US/10/823,197
; CURRENT FILING DATE: 2004-04-12
; PRIOR APPLICATION NUMBER: 09/578,534
; PRIOR FILING DATE: 2000-05-24
; PRIOR APPLICATION NUMBER: 09/431,352
; PRIOR FILING DATE: 1999-11-01
; NUMBER OF SEQ ID NOS: 29
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 7
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Primer
US-10-823-197-7

Query Match 100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 14;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
| | | | | | | | | | | | | | | | |
Db 1 CTCAGCAACACTCCTAT 17

RESULT 8
US-10-712-882-5
; Sequence 5, Application US/10712882
; Publication No. US20050032077A1
; GENERAL INFORMATION:
; APPLICANT: Duff, Gordon W.
; APPLICANT: Richardson, Robert R.S.
; APPLICANT: Rennie, Ian G.
; TITLE OF INVENTION: DETECTING GENETIC PREDISPOSITION TO
; TITLE OF INVENTION: SIGHT-THREATENING DIABETIC RETINOPATHY
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: FOLEY, HOAG & ELIOT LLP
; STREET: One Post Office Square
; CITY: Boston
; STATE: MA
; COUNTRY: USA
; ZIP: 02109-2170
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS


```
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
;   APPLICATION NUMBER: US/10/712,882
;   FILING DATE: 12-NOV-2003
;   CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
;   APPLICATION NUMBER: US/09/037,472
;   FILING DATE: 10-MAR-1998
;   APPLICATION NUMBER: PCT/GB97/02790
;   FILING DATE: 09-OCT-1997
; ATTORNEY/AGENT INFORMATION:
;   NAME: Arnold, Beth E.
;   REGISTRATION NUMBER: 35,430
; TELECOMMUNICATION INFORMATION:
;   TELEPHONE: (617) 832-1000
;   TELEFAX: (617) 832-7000
; INFORMATION FOR SEQ ID NO: 5:
;   SEQUENCE CHARACTERISTICS:
;     LENGTH: 17 base pairs
;     TYPE: nucleic acid
;     STRANDEDNESS: single
;     TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; DESCRIPTION: /desc = "primer"
; SEQUENCE DESCRIPTION: SEQ ID NO: 5:
US-10-712-882-5

Query Match      100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 14;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 9
US-10-838-503-9
; Sequence 9, Application US/10838503
; Publication No. US2005064453A1
; GENERAL INFORMATION:
; APPLICANT: DUFF, GORDON
; APPLICANT: KORNMAN, KENNETH
; APPLICANT: VAN DIJK, SIMON
; TITLE OF INVENTION: DIAGNOSTICS AND THERAPEUTICS FOR EARLY-ONSET MENOPAUSE
; FILE REFERENCE: MSA-012.01
; CURRENT APPLICATION NUMBER: US/10/838,503
; CURRENT FILING DATE: 2004-05-03
; NUMBER OF SEQ ID NOS: 30
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 9
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic
; OTHER INFORMATION: oligonucleotide
US-10-838-503-9

Query Match      100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 14;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 10
US-11-121-634-22
; Sequence 22, Application US/11121634
; Publication No. US20050282198A1
; GENERAL INFORMATION:
```

```
; APPLICANT: Duff, Gordon
; TITLE OF INVENTION: Diagnostics and Therapeutics for Diseases Associated with an IL-1
; FILE REFERENCE: 24299-508CON3 CIP1
; CURRENT APPLICATION NUMBER: US/11/121,634
; CURRENT FILING DATE: 2005-05-03
; PRIOR APPLICATION NUMBER: US 10/802,061
; PRIOR FILING DATE: 2004-03-15
; PRIOR APPLICATION NUMBER: US 09/845,129
; PRIOR FILING DATE: 2001-04-27
; PRIOR APPLICATION NUMBER: US 09/345,217
; PRIOR FILING DATE: 1999-06-30
; PRIOR APPLICATION NUMBER: PCT/GB98/01481
; PRIOR FILING DATE: 1998-05-21
; PRIOR APPLICATION NUMBER: GB9711040.7
; PRIOR FILING DATE: 1997-05-29
; PRIOR APPLICATION NUMBER: US 10/300,011
; PRIOR FILING DATE: 2002-11-19
; PRIOR APPLICATION NUMBER: US 60/386,020
; PRIOR FILING DATE: 2002-06-05
; PRIOR APPLICATION NUMBER: US 60/331,681
; PRIOR FILING DATE: 2001-11-19
; PRIOR APPLICATION NUMBER: US 60/567,727
; PRIOR FILING DATE: 2004-05-03
; NUMBER OF SEQ ID NOS: 202
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 22
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: chemically synthesized primer
US-11-121-634-22

Query Match      100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 14;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 11
US-10-626-830-9
; Sequence 9, Application US/10626830
; Publication No. US2004022928A1
; GENERAL INFORMATION:
; APPLICANT: Sims, John E.
; APPLICANT: Kaufman, Dixon B.
; TITLE OF INVENTION: IL-1 Genotype in Early Kidney Allograft Rejection
; FILE REFERENCE: WESTERN-08309
; CURRENT APPLICATION NUMBER: US/10/626,830
; CURRENT FILING DATE: 2003-07-24
; NUMBER OF SEQ ID NOS: 18
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 9
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic
US-10-626-830-9

Query Match      82.4%; Score 14; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 35;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCC 14
Db 4 CTCAGCAACACTCC 17
```

RESULT 12

US-11-083-784-1215680/c
; Sequence 1215680, Application US/11083784
; Publication No. US20050245475A1
; GENERAL INFORMATION:
; APPLICANT: Dharmacon, Inc.
; APPLICANT: Khvorova, Anastasia
; APPLICANT: Reynolds, Angela
; APPLICANT: Leake, Devin
; APPLICANT: Marshall, William
; APPLICANT: Scaringe, Stephen
; TITLE OF INVENTION: Functional and Hyperfunctional siRNA
; FILE REFERENCE: 13499US
; CURRENT APPLICATION NUMBER: US/11/083,784
; CURRENT FILING DATE: 2005-03-18
; PRIOR FILING DATE: US/10/714,333
; PRIOR FILING DATE: 2003-11-14
; PRIOR APPLICATION NUMBER: 60/502,050
; PRIOR FILING DATE: 2003-09-10
; PRIOR APPLICATION NUMBER: 60/426,137
; PRIOR FILING DATE: 2002-11-14
; NUMBER OF SEQ ID NOS: 1591911
; SOFTWARE: Proprietary
; SEQ ID NO 1215680
; TYPE: RNA
; ORGANISM: Homo sapiens
US-11-083-784-1215680

Query Match 82.4%; Score 14; DB 1; Length 19;
Best Local Similarity 100.0%; Pred. No. 39;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 CAGCAACACTCCTA 16
|||||

Db 16 CAGCAACACTCCTA 3

RESULT 13

US-11-083-784-1215694/c
; Sequence 1215694, Application US/11083784
; Publication No. US20050245475A1
; GENERAL INFORMATION:
; APPLICANT: Dharmacon, Inc.
; APPLICANT: Khvorova, Anastasia
; APPLICANT: Reynolds, Angela
; APPLICANT: Leake, Devin
; APPLICANT: Marshall, William
; APPLICANT: Scaringe, Stephen
; TITLE OF INVENTION: Functional and Hyperfunctional siRNA
; FILE REFERENCE: 13499US
; CURRENT APPLICATION NUMBER: US/11/083,784
; CURRENT FILING DATE: 2005-03-18
; PRIOR FILING DATE: US/10/714,333
; PRIOR FILING DATE: 2003-11-14
; PRIOR APPLICATION NUMBER: 60/502,050
; PRIOR FILING DATE: 2003-09-10
; PRIOR APPLICATION NUMBER: 60/426,137
; PRIOR FILING DATE: 2002-11-14
; NUMBER OF SEQ ID NOS: 1591911
; SOFTWARE: Proprietary
; SEQ ID NO 1215694
; LENGTH: 19
; TYPE: RNA
; ORGANISM: Homo sapiens
US-11-083-784-1215694

Query Match 82.4%; Score 14; DB 1; Length 19;
Best Local Similarity 100.0%; Pred. No. 39;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 CAGCAACACTCCTA 16
|||||

Db 15 CAGCAACACTCCTA 2

RESULT 14

US-11-083-784-1215717/c
; Sequence 1215717, Application US/11083784
; Publication No. US20050245475A1
; GENERAL INFORMATION:
; APPLICANT: Dharmacon, Inc.
; APPLICANT: Khvorova, Anastasia
; APPLICANT: Reynolds, Angela
; APPLICANT: Leake, Devin
; APPLICANT: Marshall, William
; APPLICANT: Scaringe, Stephen
; TITLE OF INVENTION: Functional and Hyperfunctional siRNA
; FILE REFERENCE: 13499US
; CURRENT APPLICATION NUMBER: US/11/083,784
; CURRENT FILING DATE: 2005-03-18
; PRIOR FILING DATE: US/10/714,333
; PRIOR FILING DATE: 2003-11-14
; PRIOR APPLICATION NUMBER: 60/502,050
; PRIOR FILING DATE: 2003-09-10
; PRIOR APPLICATION NUMBER: 60/426,137
; PRIOR FILING DATE: 2002-11-14
; NUMBER OF SEQ ID NOS: 1591911
; SOFTWARE: Proprietary
; SEQ ID NO 1215717
; LENGTH: 19
; TYPE: RNA
; ORGANISM: Homo sapiens
US-11-083-784-1215717

Query Match 82.4%; Score 14; DB 1; Length 19;
Best Local Similarity 100.0%; Pred. No. 39;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 CAGCAACACTCCTA 16
|||||

Db 17 CAGCAACACTCCTA 4

RESULT 15

US-11-101-244-1215680/c
; Sequence 1215680, Application US/1101244
; Publication No. US20050246794A1
; GENERAL INFORMATION:
; APPLICANT: Dharmacon, Inc.
; APPLICANT: Khvorova, Anastasia
; APPLICANT: Reynolds, Angela
; APPLICANT: Leake, Devin
; APPLICANT: Marshall, William
; APPLICANT: Scaringe, Stephen
; TITLE OF INVENTION: Functional and Hyperfunctional siRNA
; FILE REFERENCE: 13499US
; CURRENT APPLICATION NUMBER: US/11/101,244
; CURRENT FILING DATE: 2005-04-07
; PRIOR FILING DATE: 2003-09-10
; PRIOR APPLICATION NUMBER: 60/502,050
; PRIOR FILING DATE: 2003-09-10
; PRIOR APPLICATION NUMBER: 60/426,137
; PRIOR FILING DATE: 2002-11-14
; NUMBER OF SEQ ID NOS: 1591911
; SOFTWARE: Proprietary
; SEQ ID NO 1215680
; LENGTH: 19
; TYPE: RNA
; ORGANISM: Homo sapiens
US-11-101-244-1215680

Query Match 82.4%; Score 14; DB 1; Length 19;
Best Local Similarity 100.0%; Pred. No. 39;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 CAGCAACACTCCTA 16

Db 16 CAGCAACTCCTA 3
|||||

RESULT 16
US-11-101-244-1215694/c
; Sequence 1215694, Application US/11101244
; Publication No. US20050246794A1
; GENERAL INFORMATION:
; APPLICANT: Dharmacon, Inc.
; APPLICANT: Khvorova, Anastasia
; APPLICANT: Reynolds, Angela
; APPLICANT: Leake, Devin
; APPLICANT: Marshall, William
; APPLICANT: Scaringe, Stephen
; TITLE OF INVENTION: Functional and Hyperfunctional siRNA
; FILE REFERENCE: 13499US
; CURRENT APPLICATION NUMBER: US/11/101,244
; CURRENT FILING DATE: 2005-04-07
; PRIOR APPLICATION NUMBER: 60/502,050
; PRIOR FILING DATE: 2003-09-10
; PRIOR APPLICATION NUMBER: 60/426,137
; PRIOR FILING DATE: 2002-11-14
; NUMBER OF SEQ ID NOS: 1591911
; SOFTWARE: Proprietary
; SEQ ID NO 1215694
; LENGTH: 19
; TYPE: RNA
; ORGANISM: Homo sapiens
US-11-101-244-1215694

Query Match 82.4%; Score 14; DB 1; Length 19;
Best Local Similarity 100.0%; Pred. No. 39;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 CAGCAACTCCTA 16
|||||

Db 15 CAGCAACTCCTA 2

RESULT 17
US-11-101-244-1215717/c
; Sequence 1215717, Application US/11101244
; Publication No. US20050246794A1
; GENERAL INFORMATION:
; APPLICANT: Dharmacon, Inc.
; APPLICANT: Khvorova, Anastasia
; APPLICANT: Reynolds, Angela
; APPLICANT: Leake, Devin
; APPLICANT: Marshall, William
; APPLICANT: Scaringe, Stephen
; TITLE OF INVENTION: Functional and Hyperfunctional siRNA
; FILE REFERENCE: 13499US
; CURRENT APPLICATION NUMBER: US/11/101,244
; CURRENT FILING DATE: 2005-04-07
; PRIOR APPLICATION NUMBER: 60/502,050
; PRIOR FILING DATE: 2003-09-10
; PRIOR APPLICATION NUMBER: 60/426,137
; PRIOR FILING DATE: 2002-11-14
; NUMBER OF SEQ ID NOS: 1591911
; SOFTWARE: Proprietary
; SEQ ID NO 1215717
; LENGTH: 19
; TYPE: RNA
; ORGANISM: Homo sapiens
US-11-101-244-1215717

Query Match 82.4%; Score 14; DB 1; Length 19;
Best Local Similarity 100.0%; Pred. No. 39;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 CAGCAACTCCTA 16
|||||

Db 17 CAGCAACTCCTA 4

RESULT 18
US-10-310-914A-273328/c
; Sequence 273328, Application US/10310914A
; Publication No. US20060003322A1
; GENERAL INFORMATION:
; APPLICANT: Bentwich, Isaac
; APPLICANT: Shiler, Kvizat
; TITLE OF INVENTION: Bioinformatically detectable group of novel regulatory genes and
; FILE REFERENCE: 06087.0200.CPUS01
; CURRENT APPLICATION NUMBER: US/10/310,914A
; CURRENT FILING DATE: 2002-12-06
; NUMBER OF SEQ ID NOS: 1388402
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 273328
; LENGTH: 19
; TYPE: RNA
; ORGANISM: Human
US-10-310-914A-273328

Query Match 81.2%; Score 13.8; DB 1; Length 19;
Best Local Similarity 88.2%; Pred. No. 42;
Matches 15; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CTCAGCAACTCCTAT 17
|

Db 19 CCCAGCAACCTCCTAT 3

RESULT 19
US-11-083-784-921461
; Sequence 921461, Application US/11083784
; Publication No. US20050245475A1
; GENERAL INFORMATION:
; APPLICANT: Dharmacon, Inc.
; APPLICANT: Khvorova, Anastasia
; APPLICANT: Reynolds, Angela
; APPLICANT: Leake, Devin
; APPLICANT: Marshall, William
; APPLICANT: Scaringe, Stephen
; TITLE OF INVENTION: Functional and Hyperfunctional siRNA
; FILE REFERENCE: 13499US
; CURRENT APPLICATION NUMBER: US/11/083,784
; CURRENT FILING DATE: 2005-03-18
; PRIOR APPLICATION NUMBER: US/10/714,333
; PRIOR FILING DATE: 2003-11-14
; PRIOR APPLICATION NUMBER: 60/502,050
; PRIOR FILING DATE: 2003-09-10
; PRIOR APPLICATION NUMBER: 60/426,137
; PRIOR FILING DATE: 2002-11-14
; NUMBER OF SEQ ID NOS: 1591911
; SOFTWARE: Proprietary
; SEQ ID NO 921461
; LENGTH: 19
; TYPE: RNA
; ORGANISM: Homo sapiens
US-11-083-784-921461

Query Match 81.2%; Score 13.8; DB 1; Length 19;
Best Local Similarity 64.7%; Pred. No. 42;
Matches 11; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CTCAGCAACTCCTAT 17
|

Db 1 CUCAGCAUCUCUUAU 17

RESULT 20
US-11-083-784-1372880/c
; Sequence 1372880, Application US/11083784

US-11-083-784-628845/c
; Sequence 628845, Application US/11083784
; Publication No. US20050245475A1
; GENERAL INFORMATION:
; APPLICANT: Dharmacon, Inc.
; APPLICANT: Khvorova, Anastasia
; APPLICANT: Reynolds, Angela
; APPLICANT: Leake, Devin
; APPLICANT: Marshall, William
; APPLICANT: Scaringe, Stephen
; TITLE OF INVENTION: Functional and Hyperfunctional siRNA
; FILE REFERENCE: 134990S
; CURRENT APPLICATION NUMBER: US/11/083,784
; CURRENT FILING DATE: 2005-03-18
; PRIOR APPLICATION NUMBER: US/10/714,333
; PRIOR FILING DATE: 2003-11-14
; PRIOR APPLICATION NUMBER: 60/502,050
; PRIOR FILING DATE: 2003-09-10
; PRIOR APPLICATION NUMBER: 60/426,137
; PRIOR FILING DATE: 2002-11-14
; NUMBER OF SEQ ID NOS: 1591911
; SOFTWARE: Proprietary
; SEQ ID NO 628845
; LENGTH: 19
; TYPE: RNA
; ORGANISM: Homo sapiens
US-11-083-784-628845

Query Match 78.8%; Score 13.4; DB 1; Length 19;
Best Local Similarity 93.3%; Pred. No. 47;
Matches 14; Conservative 0; Mismatches 0; Indels 1; Gaps 0;

Qy 1 CTCAGCAACACTCCT 15
Db 18 CTCAGAAACACTCCT 4

RESULT 25
US-11-101-244-628637/c
; Sequence 628637, Application US/1101244
; Publication No. US20050246794A1
; GENERAL INFORMATION:
; APPLICANT: Dharmacon, Inc.
; APPLICANT: Khvorova, Anastasia
; APPLICANT: Reynolds, Angela
; APPLICANT: Leake, Devin
; APPLICANT: Marshall, William
; APPLICANT: Scaringe, Stephen
; TITLE OF INVENTION: Functional and Hyperfunctional siRNA
; FILE REFERENCE: 134990S
; CURRENT APPLICATION NUMBER: US/11/101,244
; CURRENT FILING DATE: 2005-04-07
; PRIOR APPLICATION NUMBER: 60/502,050
; PRIOR FILING DATE: 2003-09-10
; PRIOR APPLICATION NUMBER: 60/426,137
; PRIOR FILING DATE: 2002-11-14
; NUMBER OF SEQ ID NOS: 1591911
; SOFTWARE: Proprietary
; SEQ ID NO 628637
; LENGTH: 19
; TYPE: RNA
; ORGANISM: Homo sapiens
US-11-101-244-628637

Query Match 78.8%; Score 13.4; DB 1; Length 19;
Best Local Similarity 93.3%; Pred. No. 47;
Matches 14; Conservative 0; Mismatches 0; Indels 1; Gaps 0;

Qy 1 CTCAGCAACACTCCT 15
Db 18 CTCAGAAACACTCCT 4

RESULT 26
US-11-101-244-628845/c
; Sequence 628845, Application US/1101244
; Publication No. US20050246794A1
; GENERAL INFORMATION:
; APPLICANT: Dharmacon, Inc.
; APPLICANT: Khvorova, Anastasia
; APPLICANT: Reynolds, Angela
; APPLICANT: Leake, Devin
; APPLICANT: Marshall, William
; APPLICANT: Scaringe, Stephen
; TITLE OF INVENTION: Functional and Hyperfunctional siRNA
; FILE REFERENCE: 134990S
; CURRENT APPLICATION NUMBER: US/11/101,244
; CURRENT FILING DATE: 2005-04-07
; PRIOR APPLICATION NUMBER: 60/502,050
; PRIOR FILING DATE: 2003-09-10
; PRIOR APPLICATION NUMBER: 60/426,137
; PRIOR FILING DATE: 2002-11-14
; NUMBER OF SEQ ID NOS: 1591911
; SOFTWARE: Proprietary
; SEQ ID NO 628845
; LENGTH: 19
; TYPE: RNA
; ORGANISM: Homo sapiens
US-11-101-244-628845

Query Match 78.8%; Score 13.4; DB 1; Length 19;
Best Local Similarity 93.3%; Pred. No. 47;
Matches 14; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCT 15
Db 18 CTCAGAAACACTCCT 4

RESULT 27
US-10-310-914A-273305/c
; Sequence 273305, Application US/10310914A
; Publication No. US20060003322A1
; GENERAL INFORMATION:
; APPLICANT: Bentwich, Isaac
; APPLICANT: Shiller, Kvuzat
; TITLE OF INVENTION: Bioinformatically detectable group of novel regulatory genes and
; FILE REFERENCE: 06087,0200.CPUS01
; CURRENT APPLICATION NUMBER: US/10/310,914A
; CURRENT FILING DATE: 2002-12-06
; NUMBER OF SEQ ID NOS: 1388402
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 273305
; LENGTH: 18
; TYPE: RNA
; ORGANISM: Human
US-10-310-914A-273305

Query Match 75.3%; Score 12.8; DB 1; Length 18;
Best Local Similarity 87.5%; Pred. No. 52;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTA 16
Db 16 CCCAGCAACCTCCTA 1

RESULT 28
US-10-096-125-14/c
; Sequence 14, Application US/10096125
; Publication No. US20030077608A1
; GENERAL INFORMATION:
; APPLICANT: Coull, James M.
; APPLICANT: Piandaca, Mark J.
; APPLICANT: Kristjansson, Mark D.

; APPLICANT: Hyldeg-Nielsen, Jens J.
; APPLICANT: Creasey, Theresa S.
; TITLE OF INVENTION: Methods, Kits And Compositions Pertaining To
; TITLE OF INVENTION: Combination Oligomers And Libraries For Their
; TITLE OF INVENTION: Preparation
; FILE REFERENCE: BP0102-US
; CURRENT APPLICATION NUMBER: US/10/096,125
; CURRENT FILING DATE: 2002-03-09
; PRIOR APPLICATION NUMBER: 60/274,547
; PRIOR FILING DATE: 2001-03-09
; NUMBER OF SEQ ID NOS: 25
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 14
; LENGTH: 16
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic
; OTHER INFORMATION: Oligonucleotide Primer
US-10-096-125-14

Query Match 72.9%; Score 12.4; DB 1; Length 16;
Best Local Similarity 92.9%; Pred. No. 51;
Matches 13; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 TCAGCAACACTCCT 15
Db 15 TCAGCAACACTCCT 2

RESULT 29

US-10-257-017B-302368/c
; Sequence 302368, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 302368
; LENGTH: 12
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide primer for the detection of SNP TSC0019966
US-10-257-017B-302368

Query Match 64.7%; Score 11; DB 1; Length 12;
Best Local Similarity 100.0%; Pred. No. 56;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 6 CAACACTCCTA 16
Db 11 CAACACTCCTA 1

RESULT 30

US-10-257-017B-71093/c
; Sequence 71093, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 71093
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0018437
US-10-257-017B-71093

; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 71093
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0018437
US-10-257-017B-71093

Query Match 64.7%; Score 11; DB 1; Length 13;
Best Local Similarity 100.0%; Pred. No. 61;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 6 CAACACTCCTA 16
Db 13 CAACACTCCTA 3

RESULT 31

US-10-257-017B-71094
; Sequence 71094, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 71094
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0018437
US-10-257-017B-71094

Query Match 64.7%; Score 11; DB 1; Length 13;
Best Local Similarity 100.0%; Pred. No. 61;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 6 CAACACTCCTA 16
Db 1 CAACACTCCTA 11

RESULT 32

US-10-257-017B-136017/c
; Sequence 136017, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 136017
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0018437
US-10-257-017B-136017

```

LENGTH: 15
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0033969
S-10-257-017B-136017

Query Match          64.7%; Score 11; DB 1; Length 13;
Best Local Similarity 100.0%; Pred. No. 61;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

y      7 AACACTCCTAT 17
       |||||
b     12 AACACTCCTAT 2

RESULT 33
S-10-257-017B-136018
Sequence 136018, Application US/10257017B
Publication NO. US20040241651A1
GENERAL INFORMATION:
APPLICANT: Alexander Olek
APPLICANT: Christian Piepenbrock
APPLICANT: Kurt Berlin
TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and
TITLE OF INVENTION: methylations
FILE REFERENCE: E01/1193/WO
CURRENT APPLICATION NUMBER: US/10/257,017B
CURRENT FILING DATE: 2002-10-07
PRIOR APPLICATION NUMBER: DE 10019173.8
PRIOR FILING DATE: 2000-04-07
NUMBER OF SEQ ID NOS: 382046
SEQ ID NO 136018
LENGTH: 13
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0033969
S-10-257-017B-136018

Query Match          64.7%; Score 11; DB 1; Length 13;
Best Local Similarity 100.0%; Pred. No. 61;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

y      7 AACACTCCTAT 17
       |||||
b     12 AACACTCCTAT 12

RESULT 34
S-09-848-754A-9312
Sequence 9312, Application US/09848754A
Publication NO. US20030073207A1
GENERAL INFORMATION:
APPLICANT: Ribozyme Pharmaceuticals, Inc.
TITLE OF INVENTION: Enzymatic Nucleic Acid Treatment of Diseases or Conditions
TITLE OF INVENTION: Levels of Epidermal Growth Factor Receptors
FILE REFERENCE: MEHB00-958-I (400/018)
CURRENT APPLICATION NUMBER: US/09/848,754A
CURRENT FILING DATE: 2001-05-03
NUMBER OF SEQ ID NOS: 9645
SOFTWARE: PatentIn version 3.0
SEQ ID NO 9312
LENGTH: 15
TYPE: RNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic
S-09-848-754A-9312

Query Match          63.5%; Score 10.8; DB 1; Length 15;
Best Local Similarity 71.4%; Pred. No. 75;
Matches 10; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

```

```
US-10-257-017B-51767/c
; Sequence 51767, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 51767
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0014432
US-10-257-017B-51767

Query Match      61.2%; Score 10.4; DB 1; Length 13;
Best Local Similarity 91.7%; Pred. No. 72;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      6 CAACACTCCTAT 17
Db      13 CAACATTCCTAT 2

RESULT 38
US-10-257-017B-51768
; Sequence 51768, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 51768
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0014432
US-10-257-017B-51768

Query Match      61.2%; Score 10.4; DB 1; Length 13;
Best Local Similarity 91.7%; Pred. No. 72;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      6 CAACACTCCTAT 17
Db      1 CAACATTCCTAT 12

RESULT 39
US-10-257-017B-159811/c
; Sequence 159811, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
```

```
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 159811
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0040225
US-10-257-017B-159811

Query Match      61.2%; Score 10.4; DB 1; Length 13;
Best Local Similarity 91.7%; Pred. No. 72;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      3 CAGCAACACTCC 14
Db      12 CAACAACTCC 1

RESULT 40
US-10-257-017B-159812
; Sequence 159812, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 159812
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0040225
US-10-257-017B-159812

Query Match      61.2%; Score 10.4; DB 1; Length 13;
Best Local Similarity 91.7%; Pred. No. 72;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      3 CAGCAACACTCC 14
Db      2 CAACAACTCC 13

RESULT 41
US-10-257-017B-163329/c
; Sequence 163329, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
```



```
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 163329
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0041059
US-10-257-017B-163329

Query Match          61.2%; Score 10.4; DB 1; Length 13;
Best Local Similarity 91.7%; Pred. No. 72;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      4 AGCAACTCTCT 15
Db      12 ACCAACTCTCT 1

RESULT 42
US-10-257-017B-163330
; Sequence 163330, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 163330
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0041059
US-10-257-017B-163330

Query Match          61.2%; Score 10.4; DB 1; Length 13;
Best Local Similarity 91.7%; Pred. No. 72;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      4 AGCAACTCTCT 15
Db      2 ACCAACTCTCT 13

RESULT 43
US-10-257-017B-179987/c
; Sequence 179987, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 179987
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0044568
```

```
US-10-257-017B-179987

Query Match          61.2%; Score 10.4; DB 1; Length 13;
Best Local Similarity 91.7%; Pred. No. 72;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      1 CTCAGCAACT 12
Db      12 CTCATCAACT 1

RESULT 44
US-10-257-017B-179988
; Sequence 179988, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 179988
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0044568
US-10-257-017B-179988

Query Match          61.2%; Score 10.4; DB 1; Length 13;
Best Local Similarity 91.7%; Pred. No. 72;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      1 CTCAGCAACT 12
Db      2 CTCATCAACT 13

RESULT 45
US-10-257-017B-191043/c
; Sequence 191043, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 191043
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0047001
US-10-257-017B-191043

Query Match          61.2%; Score 10.4; DB 1; Length 13;
Best Local Similarity 91.7%; Pred. No. 72;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      6 CAACACTCTAT 17
```

Db 12 CACCACCTCCTAT 1
||| ||||| |||||

RESULT 46

US-10-257-017B-191044
; Sequence 191044, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; PRIOR FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 191044
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0047001
US-10-257-017B-191044

Query Match 61.2%; Score 10.4; DB 1; Length 13;
Best Local Similarity 91.7%; Pred. No. 72;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 6 CACCACCTCCTAT 17
||| ||||| |||||
Db 2 CACCACCTCCTAT 13

RESULT 47

US-10-257-017B-222887/c
; Sequence 222887, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; PRIOR FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 222887
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0006573
US-10-257-017B-222887

Query Match 61.2%; Score 10.4; DB 1; Length 13;
Best Local Similarity 91.7%; Pred. No. 72;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 6 CACCACCTCCTAT 17
||| ||||| |||||
Db 13 CAAAACCTCCTAT 2

RESULT 48

US-10-257-017B-222888
; Sequence 222888, Application US/10257017B

; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; PRIOR FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 222888
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0006573
US-10-257-017B-222888

Query Match 61.2%; Score 10.4; DB 1; Length 13;
Best Local Similarity 91.7%; Pred. No. 72;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 6 CACCACCTCCTAT 17
||| ||||| |||||
Db 1 CAAAACCTCCTAT 12

RESULT 49

US-10-257-017B-223803/c
; Sequence 223803, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 223803
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0054492
US-10-257-017B-223803

Query Match 61.2%; Score 10.4; DB 1; Length 13;
Best Local Similarity 91.7%; Pred. No. 72;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 6 CACCACCTCCTAT 17
||| ||||| |||||
Db 13 CAAAACCTCCTAT 2

RESULT 50

US-10-257-017B-223804
; Sequence 223804, Application US/10257017B
; Publication No. US20040241651A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Detection of single nucleotide polymorphisms [SNPs] and cytosine
; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 223804
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0054492
US-10-257-017B-223804

; FILE REFERENCE: E01/1193/WO
; CURRENT APPLICATION NUMBER: US/10/257,017B
; CURRENT FILING DATE: 2002-10-07
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 382046
; SEQ ID NO 223804
; LENGTH: 13
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide for detection of SNP TSC0054492
US-10-257-017B-223804

Query Match 61.2%; Score 10.4; DB 1; Length 13;
Best Local Similarity 91.7%; Pred. No. 72;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 6 CAACACTCCTAT 17
||| |||
Db 1 CAACATTCCTAT 12

Search completed: November 21, 2006, 14:32:49
Job time : 0.001 secs

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GenCore version 5.1.9
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OM nucleic - nucleic search, using sw model

Run on: November 21, 2006, 14:35:21 ; Search time 0.001 seconds
(without alignment)
0.986 Million cell updates/sec

Title: US-10-823-197-7

Perfect score: 17

Sequence: 1 CTCAGCAACTCCTAT 17

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 0.5

Searched: 3 seqs, 29 residues

Total number of hits satisfying chosen parameters: 6

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 3 summaries

Database : rstdb:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result	No.	Score	Query Match	Length	ID	Description
CF543243	1	8.4	49.4	12	1	ACCESSION:CF543243
C	2	6.4	37.6	9	1	ACCESSION:DR026242
C	3	6	35.3	8	1	ACCESSION:CL659535

ALIGNMENTS

RESULT 1
LOCUS CF543243 12 bp mRNA linear EST 22-SEP-2003
DEFINITION S014680-024-030-N02-SP6 MP1Z-ADIS-024-leaf Beta vulgaris cDNA clone 024-030-N02 5-PRIME, mRNA sequence.
ACCESSION CF543243
VERSION CF543243.1 GI:34891683
KEYWORDS EST.
SOURCE Beta vulgaris
ORGANISM Beta vulgaris
REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicotyledons; Caryophyllales; Amaranthaceae; Beta.
AUTHORS 1 (bases 1 to 12)
Herwig,R.; Schulz,B.; Weishaar,B.; Hennig,S.; Steinfath,M.; Drungowski,M.; Stahl,D.; Wruck,W.; Menze,A.; O'Brien,J.; Lehrach,H. and Radelof,U.
TITLE Construction of a 'unigene' cDNA clone set by oligonucleotide fingerprinting allows access to 25 000 potential sugar beet genes
JOURNAL Plant J. 32 (5), 845-857 (2002)
PUBMED 12472698
COMMENT Contact: Weishaar B
ADIS DNA core facility at MP1Z
Max-Planck-Institute for Plant Breeding Research
Carl-von-Linne Weg 10, 50829 Koeln, Germany

Fax: 00492215062851
Email: weishaa@mpiz-koeln.mpg.de
Insert Length: 12 Std Error: 0.00
Plate: 30 Row: N column: 02
Seq primer: SP6.

FEATURES

source

Location/Qualifiers

1..12
/organism="Beta vulgaris"
/mol_type="mRNA"
/cultivar="KWS2320 (double haploid, monogerm breeding line)"
/db_xref="GABI:936512"
/db_xref="taxon:161934"
/clone="024-030-N02"
/tissue_type="leaf"
/lab_host="EMDH10B"
/note="Vector: PCWFSORT6; Site1: Sali; Site 2: NotI; cDNA library from sugar beet, library provided by KWS Kleinwanzlebener Saatucht AG Einbeck, Germany, contact: b.schulz@kws.de; cloning sites Sali-NotI, primer sites and orientation:
SP6-Sali-CCAGCGTCG-5prime-cDNA-polyA-CC-NotI-T7; Note: Sequencing granted in the context of the GABI-Beet project, local PI: Dr. Katharina Schneider, coordinator: Prof. Christian Jung; Sequence submission managed by RZPD/GABI-Primary database:http://gabi.rzpd.de"

Query Match 49.4%; Score 8.4; DB 1; Length 12;

Best Local Similarity 90.0%; Pred. No. 0;

Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CTCAGCAACA 10

Dd 2 CTCAGCAACA 11

RESULT 2

DR026242/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

9 bp mRNA linear EST 26-MAY-2005
Osmo0110 F. cylindrus osmotic stress library Fragilariopsis
cylindrus cDNA clone Fcylesta35b12.s1, mRNA sequence.

DR026242

DR026242.1 GI:66748605

EST

Fragilariopsis cylindrus

Fragilariopsis cylindrus

Eukaryota; stramenopiles; Bacillariophyta; Bacillariophyceae; Bacillariophycidae; Bacillariales; Bacillariaceae; Fragilariopsis.

1 (bases 1 to 9)

Krell,A. and Gloeckner,G.

Analysis of an osmotic stress induced cDNA library of the

psychrophilic diatom Fragilariopsis cylindrus

Unpublished (2004)

Contact: Krell, Andreas; Gloeckner, Gernot

Biological Oceanography, Sea ice research; Genome Analysis

Alfred-Wegner-Institute for Polar and Marine Research; Institute

for Molecular Biotechnology

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Tel: ++49 471 48311812; ++49 3641 656440

Fax: ++49 471 48311425; ++49 3641 656255

Email: akrell@awi-bremerhaven.de; gernot@imb-jena.de

PCR Primers

FORWARD: 5'M13

BACKWARD: 3'M13

Seq primer: 5'GTAAACGACGGCCAG 3'.

FEATURES

source

Location/Qualifiers

1..9

/organism="Fragilariopsis cylindrus"

/mol_type="mRNA"

/db_xref="taxon:186039"

/clone="Fcylesta35b12.s1"

Search completed: November 21, 2006, 14:35:21
Job time : 0.001 secs

GenCore version 5.1.9
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OM nucleic - nucleic search, using sw model

Run on: November 21, 2006, 14:30:59 ; Search time 0.001 Seconds
(without alignments)
24.276 Million cell updates/sec

Title: US-10-823-197-7
Perfect score: 17
Sequence: 1 CTCGACCACTCTAT 17

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 0.5

Searched: 64 seqs, 714 residues

Total number of hits satisfying chosen parameters: 128

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 64 summaries

Database : rn1db:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query	Score	Match	Length	ID	Description
1	17	100.0	17	1	US-08-628-282-1	Sequence 1, Appli
2	17	100.0	17	1	US-08-587-911-1	Sequence 1, Appli
3	17	100.0	17	1	US-08-813-456-1	Sequence 1, Appli
4	17	100.0	17	1	US-09-345-217-22	Sequence 22, Appli
5	17	100.0	17	1	US-09-431-352-1	Sequence 1, Appli
6	17	100.0	17	1	US-09-845-129-22	Sequence 22, Appli
7	17	100.0	17	1	US-09-037-472-5	Sequence 5, Appli
8	17	100.0	17	1	US-09-578-534-7	Sequence 7, Appli
9	17	100.0	17	1	US-09-632-657-9	Sequence 9, Appli
10	17	100.0	17	1	US-09-693-555A-15	Sequence 15, Appli
11	17	100.0	17	1	US-09-584-950-13	Sequence 13, Appli
12	10.8	63.5	15	1	US-08-373-124A-95	Sequence 95, Appli
13	10.8	63.5	15	1	US-08-435-628-95	Sequence 95, Appli
14	9	52.9	11	1	US-09-249-155A-146	Sequence 146, Appli
15	9	52.9	12	1	US-09-491-356C-10	Sequence 10, Appli
16	8.4	49.4	12	1	US-09-281-418-146	Sequence 146, Appli
17	8.4	49.4	12	1	US-09-210-952-39	Sequence 39, Appli
18	8	47.1	9	1	US-09-990-186-2463	Sequence 2463, Ap
19	8	47.1	9	1	US-09-990-186-2463	Sequence 2464, Ap
20	8	47.1	10	1	US-09-508-753B-65	Sequence 65, Appli
21	8	47.1	10	1	US-09-508-753B-65	Sequence 78, Appli
22	8	47.1	11	1	US-08-891-789B-11	Sequence 11, Appli
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24	7.4	43.5	9	1	US-09-990-186-60	Sequence 60, Appli
25	7.4	43.5	9	1	US-09-990-186-61	Sequence 61, Appli
26	7.4	43.5	9	1	US-09-990-186-119	Sequence 119, Appli
27	7.4	43.5	9	1	US-09-990-186-2353	Sequence 2353, Ap
28	7.4	43.5	9	1	US-09-990-186-2356	Sequence 2356, Ap
29	7.4	43.5	9	1	US-09-990-186-2357	Sequence 2357, Ap
30	7.4	43.5	9	1	US-09-990-186-2431	Sequence 2431, Ap
31	7.4	43.5	9	1	US-09-990-186-2462	Sequence 2462, Ap
32	7.4	43.5	10	1	US-08-388-353-534	Sequence 534, Appli
33	7.4	43.5	10	1	US-08-388-353-535	Sequence 535, Appli

C 34	7.4	43.5	10	1	US-08-488-551B-534	Sequence 534, App
C 35	7.4	43.5	10	1	US-08-488-551B-535	Sequence 535, App
C 36	7.4	43.5	10	1	US-08-088-661F-37	Sequence 37, Appli
C 37	7.4	43.5	10	1	US-09-173-936B-1	Sequence 1, Appli
C 38	7.4	43.5	10	1	US-08-108-591B-35	Sequence 35, Appli
C 39	7.4	43.5	10	1	US-09-632-538C-7	Sequence 7, Appli
C 40	7.4	43.5	10	1	US-08-479-660-16	Sequence 16, Appli
C 41	7.4	43.5	10	1	US-09-627-536-7	Sequence 7, Appli
C 42	7.4	43.5	10	1	US-08-894-454-120	Sequence 120, App
C 43	7.4	43.5	10	1	US-09-525-906-10	Sequence 10, Appli
C 44	7.4	43.5	10	1	US-08-468-719A-35	Sequence 35, Appli
C 45	7.4	43.5	10	1	US-08-462-977B-35	Sequence 35, Appli
C 46	7.4	43.5	10	1	US-09-641-540-7	Sequence 7, Appli
C 47	7.4	43.5	10	1	US-09-748-710-24	Sequence 24, Appli
C 48	7.4	43.5	10	1	PCT-US96-09383-16	Sequence 16, Appli
C 49	7	41.2	8	1	US-08-859-954-7	Sequence 7, Appli
C 50	7	41.2	8	1	US-08-859-954-314	Sequence 314, App
C 51	7	41.2	8	1	US-08-859-954-335	Sequence 335, App
C 52	7	41.2	8	1	US-08-859-954-452	Sequence 452, App
C 53	7	41.2	9	1	US-09-990-186-454	Sequence 454, App
C 54	7	41.2	9	1	US-09-990-186-454	Sequence 454, App
C 55	7	41.2	10	1	US-09-593-323-55	Sequence 55, Appli
C 56	7	41.2	10	1	US-09-594-108-55	Sequence 55, Appli
C 57	7	41.2	10	1	US-09-344-300-55	Sequence 55, Appli
C 58	7	41.2	10	1	US-09-101-272G-91	Sequence 91, Appli
C 59	7	41.2	10	1	US-09-508-753B-61	Sequence 61, Appli
C 60	7	41.2	10	1	US-09-508-753B-159	Sequence 159, App
C 61	7	41.2	10	1	US-09-825-770-1	Sequence 1, Appli
C 62	7	41.2	10	1	US-10-677-496-1	Sequence 1, Appli
C 63	7	41.2	10	1	US-09-263-790-23	Sequence 23, Appli
C 64	7	41.2	10	1	US-09-721-777-3	Sequence 3, Appli

ALIGNMENTS

RESULT 1
US-08-628-282-1
; Sequence 1, Application US/08628282
; Patent No. 5698399
; GENERAL INFORMATION:
; APPLICANT: Duff, Gordon W.
; APPLICANT: Russell, Graham
; APPLICANT: Eastell, Richard
; TITLE OF INVENTION: DETECTING GENETIC PREDISPOSITION FOR
; TITLE OF INVENTION: OSTEOPOROSIS
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Aquilino & Welsh
; STREET: 2121 Crystal Drive, Suite 503
; CITY: Arlington
; STATE: Virginia
; COUNTRY: US
; ZIP: 22202
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/628,282
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Welsh, John L.
; REGISTRATION NUMBER: 33,621
; REFERENCE/DOCKET NUMBER: MSS-005
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (703) 920-1122
; TELEFAX: (703) 920-3399
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 17 base pairs

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; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-628-282-1

Query Match      100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 0.083;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 2
US-08-587-911-1
; Sequence 1, Application US/08587911
; Patent No. 5942390
; GENERAL INFORMATION:
; APPLICANT: Cominelli M.D., Fabio
; APPLICANT: Pizarro Ph.D., Theresa
; APPLICANT: Rotter M.D., Jerome I.
; APPLICANT: Yang M.D., Huiying
; TITLE OF INVENTION: METHODS OF SCREENING FOR ULCERATIVE
; TITLE OF INVENTION: COLITIS BY DETECTING AN INTERLEUKIN-1 RECEPTOR ANTAGONIST
; TITLE OF INVENTION: POLYMORPHISM
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pretty, Schroeder, Brueggemann & Clark
; STREET: 444 South Flower Street, Suite 2000
; CITY: Los Angeles
; STATE: California
; COUNTRY: U.S.A.
; ZIP: 90071
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/587,911
; FILING DATE: January 12, 1996
; CLASSIFICATION: 538
; ATTORNEY/AGENT INFORMATION:
; NAME: Whiteford Esq., Wendy A
; REGISTRATION NUMBER: 36,964
; REFERENCE/DOCKET NUMBER: P07 34246
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 213-622-7700
; TELEFAX: 213-489-4210
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 17 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
US-08-587-911-1

Query Match      100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 0.083;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 3
US-08-813-456-1
; Sequence 1, Application US/08813456
; Patent No. 6210877

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; GENERAL INFORMATION:
; APPLICANT: Francis, Sheila
; APPLICANT: Crossman, David
; APPLICANT: Duff, Gordon
; TITLE OF INVENTION: Prediction of Coronary Artery
; TITLE OF INVENTION: Disease
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Jenkins & Gilchrist
; STREET: 1100 Louisiana, Suite 1800
; CITY: Houston
; STATE: TX
; COUNTRY: USA
; ZIP: 77002
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/813,456
; FILING DATE: 10-MAR-1997
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Turley, Charles P
; REGISTRATION NUMBER: 35,723
; REFERENCE/DOCKET NUMBER: 33174-00003
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (713)9513310
; TELEFAX: (713)9513314
; TELEX:
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 17 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
US-08-813-456-1

Query Match      100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 0.083;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 4
US-09-345-217-22
; Sequence 22, Application US/09345217
; Patent No. 6268142
; GENERAL INFORMATION:
; APPLICANT: DUFF, GORDON W.
; APPLICANT: COX, ANGELA
; APPLICANT: CAMP, NICOLA J.
; APPLICANT: DIGIOVINE, FRANCESCO S.
; TITLE OF INVENTION: DIAGNOSTICS AND THERAPEUTICS FOR DISEASES ASSOCIATED
; TITLE OF INVENTION: WITH AN IL-1 INFLAMMATORY HAPLOTYPE
; FILE REFERENCE: MSA-010.02
; CURRENT APPLICATION NUMBER: US/09/345,217
; CURRENT FILING DATE: 1999-06-30
; EARLIER APPLICATION NUMBER: PCT/GB98/01481
; EARLIER FILING DATE: 1998-05-21
; EARLIER APPLICATION NUMBER: 9711040.7
; EARLIER FILING DATE: 1997-05-29
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 22

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; LENGTH: 17
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: primer
US-09-345-217-22

Query Match          100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 0.083;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 5
US-09-431-352-1
; Sequence 1, Application US/09431352
; Patent No. 6524795
; GENERAL INFORMATION:
; APPLICANT: Francis, Sheila E.
; APPLICANT: Crossman, David C.
; APPLICANT: Duff, Gordon W.
; APPLICANT: Kornman, Kenneth S.
; APPLICANT: Stephenson, Katherine
; TITLE OF INVENTION: DIAGNOSTICS AND THERAPEUTICS FOR CARDIOVASCULAR
; TITLE OF INVENTION: DISORDERS
; FILE REFERENCE: MSA-006.03
; CURRENT APPLICATION NUMBER: US/09/431,352
; CURRENT FILING DATE: 1999-11-01
; PRIOR APPLICATION NUMBER: 09/320,395
; PRIOR FILING DATE: 1999-05-26
; NUMBER OF SEQ ID NOS: 18
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: primer
US-09-431-352-1

Query Match          100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 0.083;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 6
US-09-845-129-22
; Sequence 22, Application US/09845129
; Patent No. 6706478
; GENERAL INFORMATION:
; APPLICANT: DUFF, GORDON W.
; APPLICANT: COX, ANGELA
; APPLICANT: CAMP, NICOLA J.
; APPLICANT: DIGIOVINE, FRANCESCO S.
; TITLE OF INVENTION: DIAGNOSTICS AND THERAPEUTICS FOR DISEASES ASSOCIATED
; TITLE OF INVENTION: WITH AN IL-1 INFLAMMATORY HAPLOTYPE
; FILE REFERENCE: MSA-010.02
; CURRENT APPLICATION NUMBER: US/09/845,129
; CURRENT FILING DATE: 2001-04-27
; PRIOR APPLICATION NUMBER: 09/345,217
; PRIOR FILING DATE: 1999-06-30
; PRIOR APPLICATION NUMBER: PCT/GB98/01481
; PRIOR FILING DATE: 1998-05-21
; PRIOR APPLICATION NUMBER: 9711040.7
; PRIOR FILING DATE: 1997-05-29
; NUMBER OF SEQ ID NOS: 32
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; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 22
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: primer
US-09-845-129-22

Query Match          100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 0.083;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 7
US-09-037-472-5
; Sequence 5, Application US/09037472
; Patent No. 6713253
; GENERAL INFORMATION:
; APPLICANT: Duff, Gordon W.
; APPLICANT: Richardson, Robert R.S.
; APPLICANT: Rennie, Ian G.
; TITLE OF INVENTION: DETECTING GENETIC PREDISPOSITION TO
; TITLE OF INVENTION: SIGHT-THREATENING DIABETIC RETINOPATHY
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: FOLEY, HOAG & ELIOT LLP
; STREET: One Post Office Square
; CITY: Boston
; STATE: MA
; COUNTRY: USA
; ZIP: 02109-2170
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/037,472
; FILING DATE: 10-MAR-1998
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/GB97/02790
; FILING DATE: 09-OCT-1997
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Arnold, Beth E.
; REGISTRATION NUMBER: 35,430
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 832-1000
; TELEFAX: (617) 832-7000
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 17 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; DESCRIPTION: /desc = "primer"
US-09-037-472-5

Query Match          100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 0.083;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17
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RESULT 8
US-09-578-534-7
; Sequence 7, Application US/09578534
; Patent No. 6720141
; GENERAL INFORMATION:
; APPLICANT: CROSSMAN, DAVID C.
; APPLICANT: DUFF, GORDON W.
; APPLICANT: FRANCIS, SHEILA E.
; APPLICANT: KORNMAN, KENNETH S.
; APPLICANT: STEPHENSON, KATHERINE
; TITLE OF INVENTION: DIAGNOSTICS AND THERAPEUTICS FOR RESTENOSIS
; FILE REFERENCE: MSA-017.02
; CURRENT APPLICATION NUMBER: US/09/578,534
; PRIOR FILING DATE: 2000-05-24
; PRIOR APPLICATION NUMBER: 09/431,352
; PRIOR FILING DATE: 1999-11-01
; NUMBER OF SEQ ID NOS: 29
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 7
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Primer
US-09-578-534-7

Query Match      100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 0.083;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 9
US-09-632-657-9
; Sequence 9, Application US/09632657
; Patent No. 6730476
; GENERAL INFORMATION:
; APPLICANT: DUFF, GORDON
; APPLICANT: KORNMAN, KENNETH
; APPLICANT: VAN DIJK, SIMON
; TITLE OF INVENTION: DIAGNOSTICS AND THERAPEUTICS FOR EARLY-ONSET MENOPAUSE
; FILE REFERENCE: MSA-012.01
; CURRENT APPLICATION NUMBER: US/09/632,657
; CURRENT FILING DATE: 2000-08-04
; NUMBER OF SEQ ID NOS: 30
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 9
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic
US-09-632-657-9

Query Match      100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 0.083;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 10
US-09-693-555A-15
; Sequence 15, Application US/09693555A
; Patent No. 6733967
; GENERAL INFORMATION:
; APPLICANT: KORNMAN, KENNETH
; APPLICANT: DUFF, GORDON
; APPLICANT: OFFENBACHER, STEVEN
; TITLE OF INVENTION: PETAL TESTING FOR PREDICTION OF LOW BIRTH WEIGHT
; FILE REFERENCE: MSA-009.01
; CURRENT APPLICATION NUMBER: US/09/693,555A
; CURRENT FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: PCT/US99/08794
; PRIOR FILING DATE: 1999-04-21
; PRIOR APPLICATION NUMBER: 60/082,487
; PRIOR FILING DATE: 1998-04-21
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 15
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Primer
US-09-693-555A-15

Query Match      100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 0.083;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 11
US-09-584-950-13
; Sequence 13, Application US/09584950
; Patent No. 6746839
; GENERAL INFORMATION:
; APPLICANT: DUFF, GORDON W.
; APPLICANT: DI GIOVINE, FRANCESCO S.
; APPLICANT: BARNES, PETER J.
; APPLICANT: LIM, SAMSON
; TITLE OF INVENTION: DIAGNOSTICS AND THERAPEUTICS FOR AN OBSTRUCTIVE AIRWAY
; FILE REFERENCE: MSA-005.02
; CURRENT APPLICATION NUMBER: US/09/584,950
; CURRENT FILING DATE: 2000-06-01
; NUMBER OF SEQ ID NOS: 28
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 13
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Primer
US-09-584-950-13

Query Match      100.0%; Score 17; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 0.083;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCAACACTCCTAT 17
Db 1 CTCAGCAACACTCCTAT 17

RESULT 12
US-08-373-124A-95/c
; Sequence 95, Application US/08373124A
; Patent No. 5646042
; GENERAL INFORMATION:
; APPLICANT: Stinchcomb, Dan T.
; APPLICANT: Draper, Kenneth
; APPLICANT: McSwiggen, James
; APPLICANT: Jarvis, Thale
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
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;; TITLE OF INVENTION: TREATMENT OF RESTENOSIS AND
;; TITLE OF INVENTION: CANCER USING RIBOZYMES
;; NUMBER OF SEQUENCES: 2627
;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: Lyon & Lyon
;; STREET: 633 West Fifth Street
;; CITY: Los Angeles
;; STATE: California
;; COUNTRY: U.S.A.
;; ZIP: 90071
;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
;; MEDIUM TYPE: storage
;; COMPUTER: IBM Compatible
;; OPERATING SYSTEM: IBM P.C. DOS 5.0
;; SOFTWARE: Word Perfect 5.1
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/08/373,124A
;; FILING DATE: January 13, 1995
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: 08/245,466
;; FILING DATE: May 18, 1994
;; APPLICATION NUMBER: 08/192,943
;; FILING DATE: February 7, 1994
;; APPLICATION NUMBER: 07/987,132
;; FILING DATE: December 7, 1992
;; APPLICATION NUMBER: 07/936,422
;; FILING DATE: August 26, 1992
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Warburg, Richard
;; REGISTRATION NUMBER: 32,327
;; REFERENCE/DOCKET NUMBER: 209/035
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: (213) 489-1600
;; TELEFAX: (213) 955-0440
;; TELEX: 67-3510
;; INFORMATION FOR SEQ ID NO: 95:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 15 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
US-08-373-124A-95

Query Match 63.5%; Score 10.8; DB 1; Length 15;
Best Local Similarity 85.7%; Pred. No. 2.3;
Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CTCAGCAACACTCC 14
Db 14 CTCAGCAACATTC 1

RESULT 13
US-08-435-628-95/c
; Sequence 95, Application US/08435628
; Patent No. 5817796
; GENERAL INFORMATION:
; APPLICANT: Stinchcomb, Dan T.
; APPLICANT: Draper, Kenneth
; APPLICANT: McSwigen, James
; APPLICANT: Jarvis, Thale
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
; TITLE OF INVENTION: TREATMENT OF RESTENOSIS AND
; TITLE OF INVENTION: CANCER USING RIBOZYMES
; NUMBER OF SEQUENCES: 2627
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Lyon & Lyon
; STREET: 633 West Fifth Street
; CITY: Los Angeles
; STATE: California

;; COUNTRY: U.S.A.
;; ZIP: 90071
;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
;; MEDIUM TYPE: storage
;; COMPUTER: IBM Compatible
;; OPERATING SYSTEM: IBM P.C. DOS 5.0
;; SOFTWARE: Word Perfect 5.1
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/08/435,628
;; FILING DATE: 05-MAY-1995
;; CLASSIFICATION: 514
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: 08/373,124
;; FILING DATE: January 13, 1995
;; APPLICATION NUMBER: 08/245,466
;; FILING DATE: May 18, 1994
;; APPLICATION NUMBER: 08/192,943
;; FILING DATE: February 7, 1994
;; APPLICATION NUMBER: 07/987,132
;; FILING DATE: December 7, 1992
;; APPLICATION NUMBER: 07/936,422
;; FILING DATE: August 26, 1992
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Warburg, Richard
;; REGISTRATION NUMBER: 32,327
;; REFERENCE/DOCKET NUMBER: 209/035
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: (213) 489-1600
;; TELEFAX: (213) 955-0440
;; TELEX: 67-3510
;; INFORMATION FOR SEQ ID NO: 95:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 15 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
US-08-435-628-95

Query Match 63.5%; Score 10.8; DB 1; Length 15;
Best Local Similarity 85.7%; Pred. No. 2.3;
Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CTCAGCAACACTCC 14
Db 14 CTCAGCAACATTC 1

RESULT 14
US-09-249-155A-146/c
; Sequence 146, Application US/09249155A
; Patent No. 6538173
; GENERAL INFORMATION:
; APPLICANT: Heber-Katz, Ellen
; TITLE OF INVENTION: Compositions and Methods for Wound
; FILE REFERENCE: 00486,78503
; CURRENT APPLICATION NUMBER: US/09/249,155A
; CURRENT FILING DATE: 1999-02-12
; PRIOR APPLICATION NUMBER: US 60/074,737
; PRIOR FILING DATE: 1998-02-13
; PRIOR APPLICATION NUMBER: US 60/097,937
; PRIOR FILING DATE: 1998-08-26
; PRIOR APPLICATION NUMBER: US 60/102,051
; PRIOR FILING DATE: 1998-09-28
; NUMBER OF SEQ ID NOS: 346
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 146
; LENGTH: 11
; TYPE: DNA
; ORGANISM: Mus musculus
US-09-249-155A-146


```
QY 7 AACACTCC 14
    |||||
Db 8 AACACTCC 1

RESULT 19
US-09-990-186-2464/c
; Sequence 2464, Application US/09990186
; Patent No. 7030215
; GENERAL INFORMATION:
; APPLICANT: LIU, Qiang
; TITLE OF INVENTION: POSITION DEPENDENT RECOGNITION OF GNN NUCLEOTIDE
; FILE REFERENCE: 8325-0011.21 / S11-US3
; CURRENT APPLICATION NUMBER: US/09/990,186
; CURRENT FILING DATE: 2001-11-20
; NUMBER OF SEQ ID NOS: 4085
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 2464
; LENGTH: 9
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: example target
US-09-990-186-2464

Query Match 47.1%; Score 8; DB 1; Length 9;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 7 AACACTCC 14
    |||||
Db 8 AACACTCC 1

RESULT 20
US-09-508-753B-65
; Sequence 65, Application US/09508753B
; Patent No. 6544736
; GENERAL INFORMATION:
; APPLICANT: Akira SHIMAMOTO
; APPLICANT: Yasuhiro FURUICHI
; APPLICANT: Yuko SHIBATA
; APPLICANT: Hiroko FUNAKI
; APPLICANT: Eiichi OHARA
; APPLICANT: Masanori WATAHIKI
; TITLE OF INVENTION: Method for Synthesizing cDNA from mRNA sample
; FILE REFERENCE: 00162/HG
; CURRENT APPLICATION NUMBER: US/09/508,753B
; CURRENT FILING DATE: 2000-06-16
; PRIOR APPLICATION NUMBER: JP 9/270324
; PRIOR FILING DATE: 1997-09-18
; NUMBER OF SEQ ID NOS: 472
; SEQ ID NO 65
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Primer
US-09-508-753B-65

Query Match 47.1%; Score 8; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 14;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 TCAGCAAC 9
    |||||
Db 1 TCAGCAAC 8

RESULT 21
US-09-508-753B-78/c
; Sequence 78, Application US/09508753B
; Patent No. 6544736
; GENERAL INFORMATION:
; APPLICANT: Akira SHIMAMOTO
; APPLICANT: Yasuhiro FURUICHI
; APPLICANT: Yuko SHIBATA
; APPLICANT: Hiroko FUNAKI
; APPLICANT: Eiichi OHARA
; APPLICANT: Masanori WATAHIKI
; TITLE OF INVENTION: Method for Synthesizing cDNA from mRNA sample
; FILE REFERENCE: 00162/HG
; CURRENT APPLICATION NUMBER: US/09/508,753B
; CURRENT FILING DATE: 2000-06-16
; PRIOR APPLICATION NUMBER: JP 9/270324
; PRIOR FILING DATE: 1997-09-18
; NUMBER OF SEQ ID NOS: 472
; SEQ ID NO 78
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Primer
US-09-508-753B-78/c

Query Match 47.1%; Score 8; DB 1; Length 11;
Best Local Similarity 100.0%; Pred. No. 13;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGCAA 8
    |||||
Db 8 CTCAGCAA 1

RESULT 22
US-08-891-789B-11
; Sequence 11, Application US/08891789B
; Patent No. 6103466
; GENERAL INFORMATION:
; APPLICANT: Grobet, Luc; Georges, Michel
; TITLE OF INVENTION: Double-Muscling in Mammals
; NUMBER OF SEQUENCES: 52
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Blake, Cassels & Graydon
; STREET: Box 25, Commerce Court West
; CITY: Toronto
; STATE: Ontario
; ZIP: M5L 1A9
; COUNTRY: Canada
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3 1/2 inch, 1.4 Mb storage
; COMPUTER: COMPAQ, IBM PC compatible
; OPERATING SYSTEM: MS-DOS 5.1
; SOFTWARE: WORD PERFECT
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/891,789B
; FILING DATE: July 14, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Hunt, John C.
; REGISTRATION NUMBER: 36,424
; REFERENCE/DOCKET NUMBER: 52836/00004
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (416) 863-4344
; TELEFAX: (416) 863-2653
; INFORMATION FOR SEQ ID NO: 11:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 11 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-891-789B-11

Query Match 47.1%; Score 8; DB 1; Length 11;
Best Local Similarity 100.0%; Pred. No. 13;
```

Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 7 AACACTCC 14
| | | | |
Db 4 AACACTCC 11

RESULT 23

US-09-990-186-59/c
; Sequence 59, Application US/09990186
; Patent No. 7030215
; GENERAL INFORMATION:
; APPLICANT: LIU, Qiang
; TITLE OF INVENTION: POSITION DEPENDENT RECOGNITION OF GNN NUCLEOTIDE
; FILE REFERENCE: 8325-0011.21 / S11-US3
; CURRENT APPLICATION NUMBER: US/09/990,186
; CURRENT FILING DATE: 2001-11-20
; NUMBER OF SEQ ID NOS: 4085
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 59
; LENGTH: 9
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: example target
; OTHER INFORMATION: DNA
US-09-990-186-59

Query Match 43.5%; Score 7.4; DB 1; Length 9;
Best Local Similarity 88.9%; Pred. No. 1.3e+02;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CTCAGCAAC 9
| | | | |
Db 9 CTCAGCATC 1

RESULT 24

US-09-990-186-60/c
; Sequence 60, Application US/09990186
; Patent No. 7030215
; GENERAL INFORMATION:
; APPLICANT: LIU, Qiang
; TITLE OF INVENTION: POSITION DEPENDENT RECOGNITION OF GNN NUCLEOTIDE
; FILE REFERENCE: 8325-0011.21 / S11-US3
; CURRENT APPLICATION NUMBER: US/09/990,186
; CURRENT FILING DATE: 2001-11-20
; NUMBER OF SEQ ID NOS: 4085
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 60
; LENGTH: 9
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: example target
; OTHER INFORMATION: DNA
US-09-990-186-60

Query Match 43.5%; Score 7.4; DB 1; Length 9;
Best Local Similarity 88.9%; Pred. No. 1.3e+02;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CTCAGCAAC 9
| | | | |
Db 9 CTCAGCATC 1

RESULT 25

US-09-990-186-61/c
; Sequence 61, Application US/09990186
; Patent No. 7030215

; GENERAL INFORMATION:
; APPLICANT: LIU, Qiang
; TITLE OF INVENTION: POSITION DEPENDENT RECOGNITION OF GNN NUCLEOTIDE
; FILE REFERENCE: 8325-0011.21 / S11-US3
; CURRENT APPLICATION NUMBER: US/09/990,186
; CURRENT FILING DATE: 2001-11-20
; NUMBER OF SEQ ID NOS: 4085
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 61
; LENGTH: 9
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: example target
; OTHER INFORMATION: DNA
US-09-990-186-61

Query Match 43.5%; Score 7.4; DB 1; Length 9;
Best Local Similarity 88.9%; Pred. No. 1.3e+02;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CTCAGCAAC 9
| | | | |
Db 9 CTCAGCATC 1

RESULT 26

US-09-990-186-119/c
; Sequence 119, Application US/09990186
; Patent No. 7030215
; GENERAL INFORMATION:
; APPLICANT: LIU, Qiang
; TITLE OF INVENTION: POSITION DEPENDENT RECOGNITION OF GNN NUCLEOTIDE
; FILE REFERENCE: 8325-0011.21 / S11-US3
; CURRENT APPLICATION NUMBER: US/09/990,186
; CURRENT FILING DATE: 2001-11-20
; NUMBER OF SEQ ID NOS: 4085
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 119
; LENGTH: 9
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: example target
; OTHER INFORMATION: DNA
US-09-990-186-119

Query Match 43.5%; Score 7.4; DB 1; Length 9;
Best Local Similarity 88.9%; Pred. No. 1.3e+02;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CTCAGCAAC 9
| | | | |
Db 9 CTCAGCATC 1

RESULT 27

US-09-990-186-2353/c
; Sequence 2353, Application US/09990186
; Patent No. 7030215
; GENERAL INFORMATION:
; APPLICANT: LIU, Qiang
; TITLE OF INVENTION: POSITION DEPENDENT RECOGNITION OF GNN NUCLEOTIDE
; FILE REFERENCE: 8325-0011.21 / S11-US3
; CURRENT APPLICATION NUMBER: US/09/990,186
; CURRENT FILING DATE: 2001-11-20
; NUMBER OF SEQ ID NOS: 4085
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 2353
; LENGTH: 9

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; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: example target
; OTHER INFORMATION: DNA
US-09-990-186-2353

Query Match      43.5%; Score 7.4; DB 1; Length 9;
Best Local Similarity 88.9%; Pred. No. 1.3e+02;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy      8 ACACCTCCTA 16
Db      9 AAACCTCCTA 1
      |||||

RESULT 28
US-09-990-186-2356/c
; Sequence 2356, Application US/09990186
; Patent No. 7030215
; GENERAL INFORMATION:
; APPLICANT: LIU, Qiang
; TITLE OF INVENTION: POSITION DEPENDENT RECOGNITION OF GNN NUCLEOTIDE
; TITLE OF INVENTION: TRIPLETS BY ZINC FINGERS
; FILE REFERENCE: 8325-0011.21 / S11-US3
; CURRENT APPLICATION NUMBER: US/09/990,186
; CURRENT FILING DATE: 2001-11-20
; NUMBER OF SEQ ID NOS: 4085
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 2356
; LENGTH: 9
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: example target
; OTHER INFORMATION: DNA
US-09-990-186-2356

Query Match      43.5%; Score 7.4; DB 1; Length 9;
Best Local Similarity 88.9%; Pred. No. 1.3e+02;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy      8 ACACCTCCTA 16
Db      9 AAACCTCCTA 1
      |||||

RESULT 29
US-09-990-186-2357/c
; Sequence 2357, Application US/09990186
; Patent No. 7030215
; GENERAL INFORMATION:
; APPLICANT: LIU, Qiang
; TITLE OF INVENTION: POSITION DEPENDENT RECOGNITION OF GNN NUCLEOTIDE
; TITLE OF INVENTION: TRIPLETS BY ZINC FINGERS
; FILE REFERENCE: 8325-0011.21 / S11-US3
; CURRENT APPLICATION NUMBER: US/09/990,186
; CURRENT FILING DATE: 2001-11-20
; NUMBER OF SEQ ID NOS: 4085
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 2357
; LENGTH: 9
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: example target
; OTHER INFORMATION: DNA
US-09-990-186-2357

Query Match      43.5%; Score 7.4; DB 1; Length 9;
Best Local Similarity 88.9%; Pred. No. 1.3e+02;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy      8 ACACCTCCTA 16
Db      9 AAACCTCCTA 1
      |||||

RESULT 29
US-09-990-186-2357/c
; Sequence 2357, Application US/09990186
; Patent No. 7030215
; GENERAL INFORMATION:
; APPLICANT: LIU, Qiang
; TITLE OF INVENTION: POSITION DEPENDENT RECOGNITION OF GNN NUCLEOTIDE
; TITLE OF INVENTION: TRIPLETS BY ZINC FINGERS
; FILE REFERENCE: 8325-0011.21 / S11-US3
; CURRENT APPLICATION NUMBER: US/09/990,186
; CURRENT FILING DATE: 2001-11-20
; NUMBER OF SEQ ID NOS: 4085
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 2357
; LENGTH: 9
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: example target
; OTHER INFORMATION: DNA
US-09-990-186-2357

Query Match      43.5%; Score 7.4; DB 1; Length 9;
Best Local Similarity 88.9%; Pred. No. 1.3e+02;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
Oy      8 ACACCTCCTA 16
Db      9 AAACCTCCTA 1
      |||||

RESULT 30
US-09-990-186-2431/c
; Sequence 2431, Application US/09990186
; Patent No. 7030215
; GENERAL INFORMATION:
; APPLICANT: LIU, Qiang
; TITLE OF INVENTION: POSITION DEPENDENT RECOGNITION OF GNN NUCLEOTIDE
; TITLE OF INVENTION: TRIPLETS BY ZINC FINGERS
; FILE REFERENCE: 8325-0011.21 / S11-US3
; CURRENT APPLICATION NUMBER: US/09/990,186
; CURRENT FILING DATE: 2001-11-20
; NUMBER OF SEQ ID NOS: 4085
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 2431
; LENGTH: 9
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: example target
; OTHER INFORMATION: DNA
US-09-990-186-2431

Query Match      43.5%; Score 7.4; DB 1; Length 9;
Best Local Similarity 88.9%; Pred. No. 1.3e+02;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy      8 ACACCTCCTA 16
Db      9 ACACCTCCCA 1
      |||||

RESULT 31
US-09-990-186-2462/c
; Sequence 2462, Application US/09990186
; Patent No. 7030215
; GENERAL INFORMATION:
; APPLICANT: LIU, Qiang
; TITLE OF INVENTION: POSITION DEPENDENT RECOGNITION OF GNN NUCLEOTIDE
; TITLE OF INVENTION: TRIPLETS BY ZINC FINGERS
; FILE REFERENCE: 8325-0011.21 / S11-US3
; CURRENT APPLICATION NUMBER: US/09/990,186
; CURRENT FILING DATE: 2001-11-20
; NUMBER OF SEQ ID NOS: 4085
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 2462
; LENGTH: 9
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: example target
; OTHER INFORMATION: DNA
US-09-990-186-2462

Query Match      43.5%; Score 7.4; DB 1; Length 9;
Best Local Similarity 88.9%; Pred. No. 1.3e+02;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy      8 ACACCTCCTA 16
Db      9 ACACCTCCCA 1
      |||||

RESULT 32
US-08-388-353-534/c
; Sequence 534, Application US/08388353
; Patent No. 6010895
; GENERAL INFORMATION:
; APPLICANT: Deacon, Nicholas J.
```

```
; APPLICANT: Learmont, Jennifer C.
; APPLICANT: McPhee, Dale A.
; APPLICANT: Crowe, Suzanne
; APPLICANT: Cooper, David
; TITLE OF INVENTION: NON-PATHOGENIC STRAINS OF HIV-1
; NUMBER OF SEQUENCES: 800
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Scully, Scott, Murphy & Presser
; STREET: 400 Garden City Plaza
; CITY: Garden City
; STATE: New York
; COUNTRY: United States
; ZIP: 11530
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA: US/08/388,353
; APPLICATION NUMBER: US/08/388,353
; FILING DATE: 14-FEB-1995
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: DiGiglio, Frank S.
; REGISTRATION NUMBER: 31,346
; REFERENCE/DOCKET NUMBER: 9606
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (516) 742-4343
; TELEFAX: (516) 742-4366
; TELEX: 230 901 SANS UR
; INFORMATION FOR SEQ ID NO: 534:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 10 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-08-388-353-534

Query Match 43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 7 AACACTCCT 15
Db 10 AACACTTCT 2
||||| |||

RESULT 33
US-08-388-353-535/c
; Sequence 535, Application US/08388353
; Patent No. 6010895
; GENERAL INFORMATION:
; APPLICANT: Deacon, Nicholas J.
; APPLICANT: Learmont, Jennifer C.
; APPLICANT: McPhee, Dale A.
; APPLICANT: Crowe, Suzanne
; APPLICANT: Cooper, David
; TITLE OF INVENTION: NON-PATHOGENIC STRAINS OF HIV-1
; NUMBER OF SEQUENCES: 800
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Scully, Scott, Murphy & Presser
; STREET: 400 Garden City Plaza
; CITY: Garden City
; STATE: New York
; COUNTRY: United States
; ZIP: 11530
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
```

```
; APPLICATION NUMBER: US/08/388,353
; FILING DATE: 14-FEB-1995
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: DiGiglio, Frank S.
; REGISTRATION NUMBER: 31,346
; REFERENCE/DOCKET NUMBER: 9606
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (516) 742-4343
; TELEFAX: (516) 742-4366
; TELEX: 230 901 SANS UR
; INFORMATION FOR SEQ ID NO: 535:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 10 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-08-388-353-535

Query Match 43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 7 AACACTCCT 15
Db 9 AACACTTCT 1
||||| |||

RESULT 34
US-08-488-551B-534/c
; Sequence 534, Application US/08488551B
; Patent No. 6015661
; GENERAL INFORMATION:
; APPLICANT: Nicholas J. Deacon
; APPLICANT: Dale A. McPhee
; APPLICANT: David Cooper
; TITLE OF INVENTION: NON-PATHOGENIC STRAINS OF HIV-1
; NUMBER OF SEQUENCES: 841
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Scully, Scott, Murphy & Presser
; STREET: 400 GARDEN CITY PLAZA
; CITY: GARDEN CITY
; STATE: NEW YORK
; COUNTRY: U.S.A.
; ZIP: 11530-0299
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/488,551B
; FILING DATE: 07-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PM3864 (AU)
; FILING DATE: 14-FEB-1994
; APPLICATION NUMBER: PM4002 (AU)
; FILING DATE: 21-FEB-1994
; APPLICATION NUMBER: PM0284 (AU)
; FILING DATE: 23-DEC-1994
; APPLICATION NUMBER: US 08/388,353
; FILING DATE: 14-FEB-1995
; APPLICATION NUMBER: PM3021/95
; FILING DATE: 17-MAY-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: FRANK S. DIGIGLIO
; REFERENCE/DOCKET NUMBER: 9606Z
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (516) 742-4343
; TELEFAX: (516) 742-4366
; INFORMATION FOR SEQ ID NO: 534:
; SEQUENCE CHARACTERISTICS:
```


LENGTH: 10 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-488-551B-534

Query Match 43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 7 AACACTCTCT 15
Db 10 AACACTTCT 2

RESULT 35
US-08-488-551B-535/c
Sequence 535, Application US/08488551B
Patent No. 6015661

GENERAL INFORMATION:
APPLICANT: Nicholas J. Deacon
APPLICANT: Dale A. McPhee
APPLICANT: David Cooper
TITLE OF INVENTION: NON-PATHOGENIC STRAINS OF HIV-1
NUMBER OF SEQUENCES: 841
CORRESPONDENCE ADDRESS:
ADDRESSEE: SCULLY, SCOTT, MURPHY & PRESSER
STREET: 400 GARDEN CITY PLAZA
CITY: GARDEN CITY
STATE: NEW YORK
COUNTRY: U.S.A.
ZIP: 11530-0299

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC Compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/488,551B
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PM3864 (AU)
FILING DATE: 14-FEB-1994
APPLICATION NUMBER: PM4002 (AU)
FILING DATE: 21-FEB-1994
APPLICATION NUMBER: PM0284 (AU)
FILING DATE: 23-DEC-1994
APPLICATION NUMBER: US 08/388,353
FILING DATE: 14-FEB-1995
APPLICATION NUMBER: PM3021/95
FILING DATE: 17-MAY-1995
ATTORNEY/AGENT INFORMATION:
NAME: FRANK S. DIGILIO
REFERENCE/DOCKET NUMBER: 9606Z
TELEPHONE: (516) 742-4343
TELEFAX: (516) 742-4366
INFORMATION FOR SEQ ID NO: 535:
SEQUENCE CHARACTERISTICS:
LENGTH: 10 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-488-551B-535

Query Match 43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 7 AACACTCTCT 15
Db 10 AACACTTCT 2

Db 9 AACACTTCT 1

RESULT 36
US-08-088-661F-37/c
Sequence 37, Application US/08088661F
Patent No. 6228982

GENERAL INFORMATION:
APPLICANT: No. 6228982den, Bengel
APPLICANT: Wittung, Pernilla
APPLICANT: Buchardt, Ole
APPLICANT: Egholm, Michael
APPLICANT: Nielsen, Peter E.
APPLICANT: Berg, Rolf
TITLE OF INVENTION: Double-Stranded Peptide Nucleic Acids
FILE REFERENCE: ISIS1108
CURRENT APPLICATION NUMBER: US/08/088,661F
CURRENT FILING DATE: 1993-07-02
PRIOR APPLICATION NUMBER: 08/054,363
PRIOR FILING DATE: 1993-04-26
PRIOR APPLICATION NUMBER: PCT/EP92/01219
PRIOR FILING DATE: 1992-05-19
NUMBER OF SEQ ID NOS: 42
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 37
LENGTH: 10
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: No. 6228982el Sequence
US-08-088-661F-37

Query Match 43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 9 CACTCTCTAT 17
Db 10 CACTACTAT 2

RESULT 37
US-09-173-936B-1/c
Sequence 1, Application US/09173936B
Patent No. 6238865

GENERAL INFORMATION:
APPLICANT: Zhen, Huang; Szostak, Jack W.
TITLE OF INVENTION: A Simple and Efficient Method to Label and Modify 3'-Terminal of RNA Using DNA Polymerase and a Synthetic Template with D Nucleotides
NUMBER OF SEQUENCES: 21
CORRESPONDENCE ADDRESS:
ADDRESSEE: Cohen, Pontani, Lieberman & Pavane
STREET: 551 Fifth Avenue
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10176
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.50 inch Diskette
COMPUTER: IBM-MS
OPERATING SYSTEM: Window 95
SOFTWARE: Microsoft Word
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/173,936B
FILING DATE: 16-Oct-1998
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/063,757
FILING DATE: 17-OCT-1997
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:

; LENGTH: 10 bases
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA
; SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-173-936B-1

Query Match 43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 3 CAGCAACAC 11
Db 9 CAGCAACCC 1

RESULT 38
US-08-108-591B-35/c
; Sequence 35, Application US/08108591B
; Patent No. 6395474
; GENERAL INFORMATION:
; APPLICANT: Buchardt, Ole
; APPLICANT: Egholm, Michael
; APPLICANT: Nielsen, Peter Bigil
; APPLICANT: Barg, Rolf Henrik
; TITLE OF INVENTION: Peptide Nucleic Acids
; FILE REFERENCE: ISIS0540
; CURRENT APPLICATION NUMBER: US/08/108,591B
; CURRENT FILING DATE: 2001-08-13
; NUMBER OF SEQ ID NOS: 43
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 35
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE: Description of Artificial Sequence
; OTHER INFORMATION: No. 6395474el Sequence
US-08-108-591B-35

Query Match 43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 9 CACTCCTAT 17
Db 10 CACTACTAT 2

RESULT 39
US-09-632-538C-7/c
; Sequence 7, Application US/09632538C
; Patent No. 6440674
; GENERAL INFORMATION:
; APPLICANT: Misra, Santosh et al.
; TITLE OF INVENTION: PLANT PROMOTER DERIVED FROM LUMINAL BINDING PROTEIN GENE AND METH
; FILE REFERENCE: 54359
; CURRENT APPLICATION NUMBER: US/09/632,538C
; CURRENT FILING DATE: 2000-08-04
; NUMBER OF SEQ ID NOS: 37
; SOFTWARE: Patentin Ver. 2.1
; SEQ ID NO 7
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: PROMOTER
; OTHER INFORMATION: ELEMENT
US-09-632-538C-7

Query Match 43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;

Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 CTCAGCAAC 9
Db 9 CTCACCAAC 1

RESULT 40
US-08-479-660-16
; Sequence 16, Application US/08479660
; Patent No. 6475806
; GENERAL INFORMATION:
; APPLICANT: Benjamin, Howard
; APPLICANT: Signer, Ethan
; APPLICANT: Geiter, Malcolm
; TITLE OF INVENTION: ANCHOR LIBRARIES AND IDENTIFICATION
; TITLE OF INVENTION: OF PEPTIDE BINDING SEQUENCES
; NUMBER OF SEQUENCES: 18
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Wolf, Greenfield & Sacks, P.C.
; STREET: 600 Atlantic Avenue
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02210
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA: US/08/479,660
; APPLICATION NUMBER: US/08/479,660
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: Greer, Helen
; REGISTRATION NUMBER: 36,816
; REFERENCE/DOCKET NUMBER: P0567/7000
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 720-3500
; TELEFAX: (617) 720-2441
; INFORMATION FOR SEQ ID NO: 16:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 10 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-479-660-16

Query Match 43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 3 CAGCAACAC 11
Db 1 CAGCAACAC 9

RESULT 41
US-09-627-536-7/c
; Sequence 7, Application US/09627536
; Patent No. 6541222
; GENERAL INFORMATION:
; APPLICANT: Misra, Santosh et al.,
; TITLE OF INVENTION: PLANT GENE PROMOTER ISOLATED FROM DOUGLAS-FIR 2S SEED STORAGE PR
; FILE REFERENCE: 54094
; CURRENT APPLICATION NUMBER: US/09/627,536
; CURRENT FILING DATE: 2000-07-28
; NUMBER OF SEQ ID NOS: 18
; SOFTWARE: Patentin Ver. 2.1
; SEQ ID NO 7
; LENGTH: 10
; TYPE: DNA

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; ORGANISM: Pseudotsuga menziesii
US-09-627-536-7

Query Match      43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      1 CTCAGCAAC 9
Db      9 CTCACCAAC 1

RESULT 42
US-08-894-454-120/c
; Sequence 120, Application US/08894454
; Patent No. 6544784
; GENERAL INFORMATION:
; APPLICANT: VAN DEN VEN, W.J.M.
; APPLICANT: SCHOENMAKERS, H.F.P.M.
; TITLE OF INVENTION: MULTIPLE-TUMOR ABERRENT GROWTH
; TITLE OF INVENTION: GENES
; NUMBER OF SEQUENCES: 164
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: The Webb Law Firm
; STREET: 700 Koppers Building, 436 Seventh Avenue
; CITY: Pittsburgh
; STATE: PA
; COUNTRY: USA
; ZIP: 15219-1818
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSEQ for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/894,454
; FILING DATE: 15-AUG-1997
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/EP/00716
; FILING DATE: 19-FEB-1996
; APPLICATION NUMBER: 95200390.3
; FILING DATE: 17-FEB-1995
; APPLICATION NUMBER: 95201951.1
; FILING DATE: 14-JUL-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Johnson, Barbara E
; REGISTRATION NUMBER: 31,198
; REFERENCE/DOCKET NUMBER: 702-971100
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 412-471-8815
; TELEFAX: 412-471-4094
; TELEX:
; INFORMATION FOR SEQ ID NO: 120:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 10 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-894-454-120

Query Match      43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      7 AACACTCCT 15
Db      10 AAAACTCCT 2

RESULT 43
US-09-525-906-10
; Sequence 10, Application US/09525906
```

```
; Patent No. 6605433
; GENERAL INFORMATION:
; APPLICANT: Jen, Jen
; APPLICANT: Sidransky, David
; APPLICANT: Kinzler, Kenneth
; APPLICANT: Vogelstein, Bert
; APPLICANT: Fliss, Makiko
; APPLICANT: Polyak, Kornelia
; TITLE OF INVENTION: Mitochondrial Dosimeter
; FILE REFERENCE: 1107.85815
; CURRENT APPLICATION NUMBER: US/09/525,906
; CURRENT FILING DATE: 2000-03-15
; PRIOR APPLICATION NUMBER: 09/377,856
; PRIOR FILING DATE: 1999-08-20
; PRIOR APPLICATION NUMBER: 60/097,307
; PRIOR FILING DATE: 1998-08-20
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 10
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-525-906-10

Query Match      43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      8 ACACCTCCTA 16
Db      2 ACACCTACTA 10

RESULT 44
US-08-468-719A-35/c
; Sequence 35, Application US/08468719A
; Patent No. 6710163
; GENERAL INFORMATION:
; APPLICANT: Buchardt, Ole
; APPLICANT: Egholm, Michael
; APPLICANT: Nielsen, Peter E.
; APPLICANT: Berg, Rolf H.
; TITLE OF INVENTION: PEPTIDE NUCLEIC ACIDS SYNTHONS
; FILE REFERENCE: ISPS-1999
; CURRENT APPLICATION NUMBER: US/08/468,719A
; CURRENT FILING DATE: 1995-06-06
; PRIOR APPLICATION NUMBER: US 08/108,591
; PRIOR FILING DATE: 1993-11-22
; NUMBER OF SEQ ID NOS: 48
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 35
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide Primer
US-08-468-719A-35

Query Match      43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      9 CACTCCTAT 17
Db      10 CACTACTAT 2

RESULT 45
US-08-462-977B-35/c
; Sequence 35, Application US/08462977B
; Patent No. 6713602
; GENERAL INFORMATION:
; APPLICANT: Buchardt, Ole
```

; APPLICANT: Egholm, Michael
; APPLICANT: Nielsen, Peter Sigil
; APPLICANT: Berg, Rolf Henrik
; TITLE OF INVENTION: Peptide Nucleic Acids
; FILE REFERENCE: IGIS-1993
; CURRENT APPLICATION NUMBER: US/08/462,977B
; CURRENT FILING DATE: 1995-06-05
; PRIOR APPLICATION NUMBER: 08/108,591
; PRIOR FILING DATE: 1993-11-22
; NUMBER OF SEQ ID NOS: 43
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 35
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY: misc feature
; OTHER INFORMATION: No. 6713602el Sequence
US-08-462-977B-35

Query Match 43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 9 CACTCTCTAT 17
Db 10 CACTACTAT 2

RESULT 46
US-09-641-540-7/c
; Sequence 7, Application US/09641540
; Patent No. 6759529
; GENERAL INFORMATION:
; APPLICANT: Miera, Santosh
; TITLE OF INVENTION: PLANT-GENE PROMOTER AND METHODS OF USING THE SAME
; FILE REFERENCE: 54358
; CURRENT APPLICATION NUMBER: US/09/641,540
; CURRENT FILING DATE: 2000-08-18
; NUMBER OF SEQ ID NOS: 28
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 7
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: PROMOTER
US-09-641-540-7

Query Match 43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 CTCAGCAAC 9
Db 9 CTCACCAAC 1

RESULT 47
US-09-748-710-24/c
; Sequence 24, Application US/09748710
; Patent No. 6916610
; GENERAL INFORMATION:
; APPLICANT: WANG, SAN MING
; APPLICANT: CHEN, JIANJUN
; APPLICANT: ROWLEY, JANET D.
; TITLE OF INVENTION: METHOD FOR GENERATION OF LONGER CDNA FRAGMENTS
; FILE REFERENCE: ARCD:343US
; CURRENT APPLICATION NUMBER: US/09/748,710
; CURRENT FILING DATE: 2000-12-22
; PRIOR APPLICATION NUMBER: 60/174,391

; PRIOR FILING DATE: 2000-01-03
; PRIOR APPLICATION NUMBER: 60/173,617
; PRIOR FILING DATE: 1999-12-29
; NUMBER OF SEQ ID NOS: 35
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 24
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic
; OTHER INFORMATION: Primer
US-09-748-710-24

Query Match 43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 TCAGCAACA 10
Db 10 TCACCAACA 2

RESULT 48
PCT-US96-09383-16
; Sequence 16, Application PC/TUS9609383
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: ANCHOR LIBRARIES AND IDENTIFICATION
; TITLE OF INVENTION: PEPTIDE BINDING SEQUENCES
; NUMBER OF SEQUENCES: 18
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Wolf, Greenfield & Sacks, P.C.
; STREET: 600 Atlantic Avenue
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02210
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US96/09383
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/479,660
; FILING DATE: 07-JUN-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Greer, Helen
; REGISTRATION NUMBER: 36,816
; REFERENCE/DOCKET NUMBER: P0567/7000WO
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 720-3500
; TELEFAX: (617) 720-2441
; INFORMATION FOR SEQ ID NO: 16:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 10 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
PCT-US96-09383-16

Query Match 43.5%; Score 7.4; DB 1; Length 10;
Best Local Similarity 88.9%; Pred. No. 19;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 3 CAGCAACAC 11
Db 1 CAGCACCAC 9

RESULT 49

US-08-859-954-7/C
; Sequence 7, Application US/08859954
; Patent No. 6083695
; GENERAL INFORMATION:
; APPLICANT: Hardin, Susan H.
; APPLICANT: Homayouni, Ramin
; APPLICANT: Hardin, Paul E.
; TITLE OF INVENTION: Design and Optimized Primer Library for
; NUMBER OF SEQUENCES: 566
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fulbright & Jaworski L.L.P.
; STREET: 1301 McKinney, Suite 5100
; CITY: Houston
; STATE: Texas
; COUNTRY: U.S.A.
; ZIP: 77010-3095

COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/859,954
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/632,782
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Paul, Thomas D.
; REGISTRATION NUMBER: 32,714
; REFERENCE/DOCKET NUMBER: D-5900
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 713/651-5325
; TELEFAX: 713/651-5246
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 8 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; DESCRIPTION: /desc = "oligonucleotide"
; HYPOTHETICAL: YES
; ANTI-SENSE: YES

US-08-859-954-7
Query Match 41.2%; Score 7; DB 1; Length 8;
Best Local Similarity 100.0%; Pred. No. 1.4e+02;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 TCAGCAA 8
Db 7 TCAGCAA 1

RESULT 50
US-08-859-954-314
; Sequence 314, Application US/08859954
; Patent No. 6083695
; GENERAL INFORMATION:
; APPLICANT: Hardin, Susan H.
; APPLICANT: Homayouni, Ramin
; APPLICANT: Hardin, Paul E.
; TITLE OF INVENTION: Design and Optimized Primer Library for
; NUMBER OF SEQUENCES: 566
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fulbright & Jaworski L.L.P.
; STREET: 1301 McKinney, Suite 5100

Query Match 41.2%; Score 7; DB 1; Length 8;
Best Local Similarity 100.0%; Pred. No. 1.4e+02;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 TCAGCAA 8
Db 7 TCAGCAA 1

RESULT 51
US-08-859-954-335/C
; Sequence 335, Application US/08859954
; Patent No. 6083695
; GENERAL INFORMATION:
; APPLICANT: Hardin, Susan H.
; APPLICANT: Homayouni, Ramin
; APPLICANT: Hardin, Paul E.
; TITLE OF INVENTION: Design and Optimized Primer Library for
; NUMBER OF SEQUENCES: 566
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fulbright & Jaworski L.L.P.
; STREET: 1301 McKinney, Suite 5100
; CITY: Houston
; STATE: Texas
; COUNTRY: U.S.A.
; ZIP: 77010-3095

COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/859,954
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/632,782

CITY: Houston
STATE: Texas
COUNTRY: U.S.A.
ZIP: 77010-3095
COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/859,954
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/632,782
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Paul, Thomas D.
; REGISTRATION NUMBER: 32,714
; REFERENCE/DOCKET NUMBER: D-5900
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 713/651-5325
; TELEFAX: 713/651-5246
; INFORMATION FOR SEQ ID NO: 314:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 8 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; DESCRIPTION: /desc = "oligonucleotide"
; HYPOTHETICAL: YES
; ANTI-SENSE: YES

US-08-859-954-314
Query Match 41.2%; Score 7; DB 1; Length 8;
Best Local Similarity 100.0%; Pred. No. 1.4e+02;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGCA 7
Db 1 CTCAGCA 7

RESULT 51
US-08-859-954-335/C
; Sequence 335, Application US/08859954
; Patent No. 6083695
; GENERAL INFORMATION:
; APPLICANT: Hardin, Susan H.
; APPLICANT: Homayouni, Ramin
; APPLICANT: Hardin, Paul E.
; TITLE OF INVENTION: Design and Optimized Primer Library for
; NUMBER OF SEQUENCES: 566
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fulbright & Jaworski L.L.P.
; STREET: 1301 McKinney, Suite 5100
; CITY: Houston
; STATE: Texas
; COUNTRY: U.S.A.
; ZIP: 77010-3095

COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/859,954
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/632,782

```
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Paul, Thomas D.
; REGISTRATION NUMBER: 32,714
; REFERENCE/DOCKET NUMBER: D-5900
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 713/651-5325
; TELEFAX: 713/651-5246
; INFORMATION FOR SEQ ID NO: 335:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 8 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; DESCRIPTION: /desc = "oligonucleotide"
; HYPOTHETICAL: YES
; ANTI-SENSE: YES
US-08-859-954-335

Query Match 41.2%; Score 7; DB 1; Length 8;
Best Local Similarity 100.0%; Pred. No. 1.4e+02;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 8 ACATCC 14
Db 8 ACATCC 2

RESULT 52
US-08-859-954-452/c
; Sequence 452, Application US/08859954
; Patent No. 6083695
; GENERAL INFORMATION:
; APPLICANT: Hardin, Susan H.
; APPLICANT: Homayouni, Ramin
; APPLICANT: Hardin, Paul E.
; TITLE OF INVENTION: Design and Optimized Primer Library for
; TITLE OF INVENTION: Gene Sequencing and Method Thereof
; NUMBER OF SEQUENCES: 566
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fulbright & Jaworski L.L.P.
; STREET: 1301 McKinney, Suite 5100
; CITY: Houston
; STATE: Texas
; COUNTRY: U.S.A.
; ZIP: 77010-3095
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/859,954
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/632,782
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Paul, Thomas D.
; REGISTRATION NUMBER: 32,714
; REFERENCE/DOCKET NUMBER: D-5900
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 713/651-5325
; TELEFAX: 713/651-5246
; INFORMATION FOR SEQ ID NO: 452:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 8 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid

; DESCRIPTION: /desc = "oligonucleotide"
; HYPOTHETICAL: YES
; ANTI-SENSE: YES
US-08-859-954-452

Query Match 41.2%; Score 7; DB 1; Length 8;
Best Local Similarity 100.0%; Pred. No. 1.4e+02;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 9 CACTCCT 15
Db 7 CACTCCT 1

RESULT 53
US-09-990-186-453/c
; Sequence 453, Application US/09990186
; Patent No. 7030215
; GENERAL INFORMATION:
; APPLICANT: LIU, Qiang
; TITLE OF INVENTION: POSITION DEPENDENT RECOGNITION OF GNN NUCLEOTIDE
; TITLE OF INVENTION: TRIPLETS BY ZINC FINGERS
; FILE REFERENCE: 8325-0011.21 / S11-US3
; CURRENT APPLICATION NUMBER: US/09/990,186
; CURRENT FILING DATE: 2001-11-20
; NUMBER OF SEQ ID NOS: 4085
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 453
; LENGTH: 9
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: example target
US-09-990-186-453

Query Match 41.2%; Score 7; DB 1; Length 9;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 CAGCAAC 9
Db 7 CAGCAAC 1

RESULT 54
US-09-990-186-454/c
; Sequence 454, Application US/09990186
; Patent No. 7030215
; GENERAL INFORMATION:
; APPLICANT: LIU, Qiang
; TITLE OF INVENTION: POSITION DEPENDENT RECOGNITION OF GNN NUCLEOTIDE
; TITLE OF INVENTION: TRIPLETS BY ZINC FINGERS
; FILE REFERENCE: 8325-0011.21 / S11-US3
; CURRENT APPLICATION NUMBER: US/09/990,186
; CURRENT FILING DATE: 2001-11-20
; NUMBER OF SEQ ID NOS: 4085
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 454
; LENGTH: 9
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: example target
US-09-990-186-454

Query Match 41.2%; Score 7; DB 1; Length 9;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 CAGCAAC 9
Db 7 CAGCAAC 1
```

```
Db          7 CAGCAAC 1

RESULT 55
US-09-593-323-55/c
; Sequence 55, Application US/09593323
; Patent No. 6265213
; GENERAL INFORMATION:
; APPLICANT: Morgan, Antony R.
; APPLICANT: Severini, Alberto
; TITLE OF INVENTION: Compositions and Methods for Determining the Activity
; TITLE OF INVENTION: of DNA-Binding Proteins and of Initiation of
; TITLE OF INVENTION: Transcription
; FILE REFERENCE: DNAB-02921
; CURRENT APPLICATION NUMBER: US/09/593,323
; CURRENT FILING DATE: 2000-06-13
; PRIOR APPLICATION NUMBER: 09/344,300
; PRIOR FILING DATE: 1999-06-24
; NUMBER OF SEQ ID NOS: 72
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 55
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic
US-09-593-323-55

Query Match          41.2%; Score 7; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy          1 CTCAGCA 7
Db          7 CTCAGCA 1

RESULT 56
US-09-594-108-55/c
; Sequence 55, Application US/09594108
; Patent No. 6284468
; GENERAL INFORMATION:
; APPLICANT: Morgan, Antony R.
; APPLICANT: Severini, Alberto
; TITLE OF INVENTION: Compositions and Methods for Determining the Activity
; TITLE OF INVENTION: of DNA-Binding Proteins and of Initiation of
; TITLE OF INVENTION: Transcription
; FILE REFERENCE: DNAB-02921
; CURRENT APPLICATION NUMBER: US/09/594,108
; CURRENT FILING DATE: 2000-06-13
; PRIOR APPLICATION NUMBER: 09/344,300
; PRIOR FILING DATE: 1999-06-24
; NUMBER OF SEQ ID NOS: 72
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 55
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic
US-09-594-108-55

Query Match          41.2%; Score 7; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy          1 CTCAGCA 7
Db          7 CTCAGCA 1

RESULT 57
US-09-344-300-55/c
; Sequence 55, Application US/09344300B
; Patent No. 6297013
; GENERAL INFORMATION:
; APPLICANT: Morgan, Antony R.
; APPLICANT: Severini, Alberto
; TITLE OF INVENTION: Compositions and Methods for Determining the Activity
; TITLE OF INVENTION: of DNA-Binding Proteins and of Initiation of
; TITLE OF INVENTION: Transcription
; FILE REFERENCE: DNAB-02921
; CURRENT APPLICATION NUMBER: US/09/344,300B
; CURRENT FILING DATE: 1999-06-24
; NUMBER OF SEQ ID NOS: 72
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 55
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic
US-09-344-300-55

Query Match          41.2%; Score 7; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy          1 CTCAGCA 7
Db          7 CTCAGCA 1

RESULT 58
US-09-101-272G-91/c
; Sequence 91, Application US/09101272G
; Patent No. 6509445
; GENERAL INFORMATION:
; APPLICANT: Nissin Food Products Co., Ltd.
; TITLE OF INVENTION: CANCEROUS METASTASIS INHIBITOR
; FILE REFERENCE: Q50979
; CURRENT APPLICATION NUMBER: US/09/101,272G
; CURRENT FILING DATE: 1998-07-08
; PRIOR APPLICATION NUMBER: JP 1059/1996
; PRIOR FILING DATE: 1996-01-08
; NUMBER OF SEQ ID NOS: 107
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 91
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; OTHER INFORMATION: 5' end of HI-8 UTI DNA fragment (coding strand)
US-09-101-272G-91

Query Match          41.2%; Score 7; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy          3 CAGCAAC 9
Db          8 CAGCAAC 2

RESULT 59
US-09-508-753B-61/c
; Sequence 61, Application US/09508753B
; Patent No. 6544736
; GENERAL INFORMATION:
; APPLICANT: Akira SHIMAMOTO
; APPLICANT: Yasuhiro FURUICHI
; APPLICANT: Yuko SHIBATA
; APPLICANT: Hiroko FUNAKI
; APPLICANT: Ei-ji OHARA
; APPLICANT: Masanori WATAHIKI
```

; TITLE OF INVENTION: Method for Synthesizing cDNA from mRNA sample
; FILE REFERENCE: 00162/HG
; CURRENT APPLICATION NUMBER: US/09/508.753B
; CURRENT FILING DATE: 2000-06-16
; PRIOR APPLICATION NUMBER: JP 9/270324
; PRIOR FILING DATE: 1997-09-18
; NUMBER OF SEQ ID NOS: 472
; SEQ ID NO 61
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Primer
US-09-508-753B-61

Query Match 41.2%; Score 7; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCA 7
| | | | |
Db 10 CTCAGCA 4

RESULT 60
US-09-508-753B-159
; Sequence 159, Application US/09508753B
; Patent No. 6544736
; GENERAL INFORMATION:
; APPLICANT: Akira SHIMAMOTO
; APPLICANT: Yasuhiro FURUICHI
; APPLICANT: Yuko SHIBATA
; APPLICANT: Hiroko FUNAKI
; APPLICANT: Eiji OHARA
; APPLICANT: Masanori WATAHIKI
; TITLE OF INVENTION: Method for Synthesizing cDNA from mRNA sample
; FILE REFERENCE: 00162/HG
; CURRENT APPLICATION NUMBER: US/09/508.753B
; CURRENT FILING DATE: 2000-06-16
; PRIOR APPLICATION NUMBER: JP 9/270324
; PRIOR FILING DATE: 1997-09-18
; NUMBER OF SEQ ID NOS: 472
; SEQ ID NO 159
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Primer
US-09-508-753B-159

Query Match 41.2%; Score 7; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCAGCA 7
| | | | |
Db 1 CTCAGCA 7

RESULT 61
US-09-825-770-1/c
; Sequence 1, Application US/09825770
; Patent No. 6686180
; GENERAL INFORMATION:
; APPLICANT: Blake, Milan S.
; APPLICANT: Bogdan, John A.
; APPLICANT: Nazario-Larrieu, Javier
; TITLE OF INVENTION: Improved Method for the Production of
; TITLE OF INVENTION: Bordatella Pertussis Toxin
; FILE REFERENCE: NV1933
; CURRENT APPLICATION NUMBER: US/09/825,770
; CURRENT FILING DATE: 2001-04-04
; PRIOR APPLICATION NUMBER: 60/194,482

; PRIOR FILING DATE: 2000-04-04
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 1
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Unknown
; FEATURE:
; OTHER INFORMATION: recombinant DNA
US-09-825-770-1

Query Match 41.2%; Score 7; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 TCAGCAA 8
| | | | |
Db 9 TCAGCAA 3

RESULT 62
US-10-677-496-1/c
; Sequence 1, Application US/10677496
; Patent No. 7018813
; GENERAL INFORMATION:
; APPLICANT: Blake, Milan S.
; APPLICANT: Bogdan, John A.
; APPLICANT: Nazario-Larrieu, Javier
; TITLE OF INVENTION: Improved Method for the Production of
; TITLE OF INVENTION: Bordatella Pertussis Toxin
; FILE REFERENCE: NV1933
; CURRENT APPLICATION NUMBER: US/10/677,496
; CURRENT FILING DATE: 2003-10-03
; PRIOR APPLICATION NUMBER: US/09/825,770
; PRIOR FILING DATE: 2001-04-04
; PRIOR APPLICATION NUMBER: 60/194,482
; PRIOR FILING DATE: 2000-04-04
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 1
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Unknown
; FEATURE:
; OTHER INFORMATION: recombinant DNA
US-10-677-496-1

Query Match 41.2%; Score 7; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 TCAGCAA 8
| | | | |
Db 9 TCAGCAA 3

RESULT 63
US-09-263-790-23
; Sequence 23, Application US/09263790
; Patent No. PPI2997
; GENERAL INFORMATION:
; APPLICANT: Nirmal Kumar PATRA et al.
; TITLE OF INVENTION: JAL PALLAVI, WATER LOGGING TOLERANT CYMBOPOGON WINTERIANUS
; FILE REFERENCE: 2761-0120P
; CURRENT APPLICATION NUMBER: US/09/263,790
; CURRENT FILING DATE: 1999-03-05
; NUMBER OF SEQ ID NOS: 38
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 23
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Artificial
; FEATURE:

Tue Nov 21 14:35:55 2006.

; OTHER INFORMATION: OPT 03 Primer - Used to develop the unique RAPD profiles of the
; OTHER INFORMATION: plant Jal Pallavi
US-09-263-790-23

Query Match 41.2%; Score 7; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 9 CACTCCT 15
| | | | |
Db 3 CACTCCT 9 .

RESULT 64

US-09-721-777-3
; Sequence 3, Application US/09721777
; Patent No. PP13279
; GENERAL INFORMATION:
; APPLICANT: Khanuja, Suman Preet Singh
; APPLICANT: Kumar, Sushil
; APPLICANT: Shasany, Ajit Kumar
; APPLICANT: Dhawan, Sunita
; APPLICANT: Darokar, Mahendra Pandurang
; APPLICANT: Naqvi, Ali Arif
; APPLICANT: Dhawan, Om Parkash
; APPLICANT: Singh, Anil Kumar
; APPLICANT: Patra, Nirmal Kumar
; APPLICANT: Bahl, Janak Raj
; APPLICANT: Bansal, Ram Prakash
; TITLE OF INVENTION: Mint Plant Named Saksham
; FILE REFERENCE: 033166-002
; CURRENT APPLICATION NUMBER: US/09/721,777
; CURRENT FILING DATE: 2000-11-27
; NUMBER OF SEQ ID NOS: 20
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 10
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: OPT primer
US-09-721-777-3

Query Match 41.2%; Score 7; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 9 CACTCCT 15
| | | | |
Db 3 CACTCCT 9

Search completed: November 21, 2006, 14:30:59
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